New Approach about the Signaling Crosstalk between IQGAPs/NF-κB/IL-8 and PDCD5/p53/TRAIL Pathways that Modulate Malignant Transformation in Hepatocellular Carcinoma

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Abstract

Objective: We aimed to investigate the signalling crosstalk of TNF-related apoptosis-inducing ligand, TRAIL death receptors, tumour protein p53, and programmed cell death (PDCD5) with IQGAPs. Also, we targeted the crosstalk between IQGAPs genes with decoy receptors, nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB), and interleukins -8 (IL-8) and its receptor genes in a designed model of hepatocellular carcinoma induced in male Balb/c mice. Methods: The presence of HCC was confirmed by histological and morphological alterations. In parallel to the incidence of hepatic cancer, we found lung, heart, and kidney cancer after treatment with DEN. Results: Our results show that the expression of mRNA of IQGAP1, TRAIL decoy receptors, NF-κB, and IL-8 genes was elevated in hepatocellular carcinoma, as compared to normal liver tissue, while their expression was further up-regulated by increasing the dose of diethylnitrosamine. The expression of IQGAP2, TRAIL death receptors, p53, and PDCD5 was significantly down-regulated in HCC (p<0.05). For confirmation of gene expression, protein levels of both IQGAP1 and P53 were measured by western blot analysis, which showed that diethylnitrosamine enhanced protein expression of IQGAP1 and diminished that of p53. Conclusion: IQGAPs have a direct signaling relationship with p53, IL-8, and TRAIL family. This interaction is recognized as a key signalling pathway for hepatocellular carcinogenesis.

Keywords: IQGAPs- IL-8- TRAIL- DEN- hepatocellular carcinoma

Introduction

N-nitrosodiethylamine (DEN) is generally utilized as a carcinogenic agent and the main organ to be targeted is species-specific. Generally, following DEN administration, mice develop not only liver cancer but also gastrointestinal, skin, respiratory and haematopoietic cancers (Saber et al., 2018; Santos et al., 2017; Zhang et al., 2019). The carcinogenicity of "DEN" is due to its ability to alkylate DNA components. It elicits its effects in a dose-dependent manner (Liu et al., 2020). A single dose of DEN with low initiation does not result in the production of carcinoma, whereas administration of a higher dose can induce Hepatocellular carcinoma (HCC) after a latency period (Zoheir et al., 2015). HCC is known to be the fifth most common solid tumour in the world, and the third most common reason of cancer-related mortality (Sayiner et al., 2019). Due to the fact that most HCC patients have acquired cirrhosis, chemotherapy regimen or major resections are not efficiently tolerated (Wang et al., 2019).

Apoptosis is a unique mode of programmed cell death that is essential for cellular homeostasis in the normal tissues and is regularly suppressed in tumorigenic processes (Jan, 2019). Deregulations of apoptosis and accelerated proliferation of hepatocytes have been declared as crucial factors, with respect to carcinogenesis of the liver or tumour progression in HCC (Czauderna et al., 2019). Basically, apoptosis occurs in cells in response to binding of ligands to their specific death receptors (DRs) through the extrinsic pathway, or to a variety of intracellular, mitochondria-initiated, stress signals which act directly on targets inside the cells in what is known as the mitochondrial or intrinsic pathway. The binding of TNF related apoptosis-inducing ligand (TRAIL) to its death receptors (DR4 and DR5) or decoy receptors (DcR1 and DcR2) is a vital example of the process regulation (Micheau, 2018). Contrary to TRAIL, NF-kB is an anti-apoptotic transcription factor that is often active in a constitutive manner in a number of cancers (Medeiros et al., 2021; Puar et al., 2018; Rasmi et al., 2020). NF-kB is actually known to induce the expression of the inflammatory CXC chemokine interleukin-8 (IL-8) in numerous cancers including HCC (Arvantakis et al., 2020).
Programmed cell death 5 (PDCD5), known also as TF-1 cell apoptosis-related gene-19 (TFAR19), was originally recognized in human leukemia TF-1 cells having serum withdrawal-induced apoptosis (Ha et al., 2021; Li et al., 2018). PDCD5 was declared downregulated in a lot of the human malignancies, including breast cancer (Hedenfalk et al., 2001) lung adenocarcinoma and hepatocellular carcinoma (Ma et al., 2019), as well as ovarian cancer (Gao et al., 2015). Accumulating evidence demonstrates that PDCD5 stimulates apoptosis of numerous human cancer cells, including TF-1, gastric cancer MGC-803, breast cancer, cervical cancer (Li et al., 2018).

We have recently shown that isoleucine-glutamine-motif containing GTPase-activating proteins (IQGAPs) have a direct and obvious signaling relation with RAS genes in production of cancer, and they are recognized as important target genes for regulation of the liver carcinogenesis (Zoheir et al., 2015). IQGAPs constitute a tiny subgroup of evolutionally preserved superfamily of GTPase-activating proteins which exist in eukaryotes ranging from yeast to the mammals. These proteins are generally abbreviated as IQGAPs due to their isoleucine-glutamine (IQ) and GTP-activating protein (GAP) domains (Xiao et al., 2018).

In this current study, we sought to investigate the signaling crosstalk of TRAIL receptors, PDCD5, p53, with IQGAPs, NF-kB, and IL-8 and its receptors in an induced model of HCC.

Materials and Methods

Ethics approval

Animals were kept in compliance with the National Institutes of Health (NIH) Guide for the Care and Use of Laboratory Animals. Medical Research Ethics Committee approved this study “Project number”12060168”.

Animals

A total of eighty 21-day-old male Balb/c mice were purchased from the Animal Care Center, and housed in polycarbonate cages in an animal facility accredited by the Association for Laboratory Animal Care Evaluation and Accreditation. All mice were fed NIH-07 diet (rodent chow) and received water ad libitum. Before the tests, the animals were allowed to acclimatize for a week. There were special training in animal care or handling provided for research staff. Every day frequently animal health and behavior were monitored to determine when animals display early markers associated with death or poor prognosis of quality of life, or specific signs of severe suffering or distress, and should be euthanized before allowing such conditions to continue or progress to death. Every day we frequently monitored the health of the animals, whether there were any unexpected deaths, and we determined cause of death in these cases.

Chemicals

DEN was bought from Sigma Chemicals (St. Louis, MO, USA). TRIZol reagent was brought from Invitrogen (Carlsbad, CA, USA). High-Capacity cDNA reverse transcription kit, primers for real time PCR, and SYBR Green Universal Master mix were all bought from Applied Biosystems® (Foster City, CA, USA). Antibodies specific for b-actin, IQGAP1 and P53 were purchased from (Santa Cruz Biotechnology).

Experimental Design

After acclimation, the 4-week-old mice were randomly divided into six experimental groups, further divided into 4-week and 8-week groups. The 4-week experimental group of mice were injected intraperitoneally (i.p.) with DEN that was dissolved in saline at the exact following dosages of 0, 50 and 75 mg/kg, groups 1-3, respectively, during the experimental days number 0, 7, 14 and 21. The i.p. injections were given in the 8-week treatment group as described above on experimental days 0, 7, 14, 21, 28, 35, 42 and 49 (groups 4-6). Mice were euthanized by CO₂ asphyxiation and exsanguination, weighed and necropsied at experimental day 288 (after injection of DEN). The mice were removed in toto, weighed and examined to find if there was any grossly visible lesion. In addition, these mice were then divided by lobe and cut into 1- to 2-mm slides. One piece from each lobe was fixed in formalin for a period of 48 to 72 hours and then was embedded in paraffin to finally give a total of three paraffin blocks per animal. Serial sections from each block were stained with hematoxylin and eosin (H&E), and hepatic preneoplastic and neoplastic lesions were classified basically on the pathology of the mouse (16).

Quantitative Analysis of Altered Hepatocellular Foci (AHF)

Quantitative analysis of AHF was achieved by operating a two-dimensional evaluation method. Whole H&E staining sections were scanned by ScanScope CS (Aperio Technologies, Vista, CA, USA). Images were then viewed with Aperio’s ImageScope viewer software (Aperio Technologies), on which the quantitative analyses were performed. The number and volume of each foci (eosinophilic, clear and basophilic) and total volume of liver sections were quantified. Because large tumors of the liver were recognized in one animal (one animal in the 75 mg/kg for 8 weeks group), these areas were not included in the total volume of liver sections of this animal. The multiplicity of AHF was reported as numbers per centimeter squared, and the volume of foci was represented as the percentage of total liver volume (%).

RNA extraction and cDNA synthesis

Under the manufacturer’s instructions, the total RNA from the liver tissue homogenate was isolated using TRIZol reagent (Gibco, Invitrogen; Eugene, OR, USA) and RNA absorbance was measured at 260 nm. The RNA quality was determined by measuring the 260/280 ratio. According to the manufacturer’s instructions, the first strand cDNA synthesis was operated by utilizing the High-Capacity cDNA reverse transcription kit (Applied Biosystems®; Foster City, CA, USA).
Quantification of mRNA expression by real-time polymerase chain reaction (RT-PCR)

To help identify the markers of apoptosis and inflammation, the Quantitative analyses of target genes mRNA expression were undertaken by RT-PCR. Briefly, the resultant cDNA from the above preparation was subjected to PCR amplification using 96-well optical reaction plates in the ABI Prism 7500 System (Applied Biosystems). The 25 μl reaction mixture contained 0.1 μl of 10 μM forward primer and 0.1 μl of 10 μM reverse primer (40 nM final concentration of each primer), 12.5 μl of SYBR Green Universal Master mix, 11.05 μl of nuclelease-free water and 1.25 μl of cDNA sample. The primers that have been used in the study were chosen from the website: pubmed.com (http://www.ncbi.nlm.nih.gov/tools/primer-blast) as listed in Table 1. To test for the contamination of any assay reagents, controls of the assay, namely, no-template controls, were incorporated onto the same plate. The real-time PCR data were analyzed utilizing the relative gene expression (i.e., ΔΔCT) method, as described in Applied Biosystems User Bulletin No. 2. The data are presented as the fold change in gene expression normalized to the endogenous reference gene glyceraldehyde-3-phosphate-dehydrogenase (GAPDH) and relative to a calibrator.

Western blot analysis

The hepatocytes were homogenized and lysed in RIPA buffer (50 mM Tris-HCl, pH 8.0, 150 mM sodium chloride, 1.0 % NP-40, 0.5% sodium deoxycholate, 0.1 % sodium dodecyl sulfate (SDS) and 0.1 % of protease inhibitor cocktail) for 20 min at 40°C. The lysates were centrifuged at 14,000 rpm at 40°C for 1.5 min to remove debris. Protein concentrations of whole cell lysates were determined using the Protein Assay Kit (Sigma Aldrich, Catalog Numbers BCA1 and B9643). To prepare 50 mg of each sample taken, the supernatant protein has been isolated by SDS-10 percent polyacrylamide gel electrophoresis (SDS-PAGE) and transferred, by standard procedures, to polyvinylide difluoride (PVDF) membrane (Bio-rad, Hercules, CA, USA). The membranes were hybridized with primary antibodies followed by incubation with appropriate secondary antibodies. By treatment with the chemi-luminescence detection reagent (Amersham Biosciences), according to manufacturer’s instructions, the antibody-bound proteins were observed after exposure to X-ray films (Kodak X-Omat). The same membranes were re-probed with anti-β actin antibody, which was used as an internal control for protein loading.

IQGAP1 and P53 antibodies as well as those rose against β-actin were diluted in PBST containing 5% non-fat milk and incubated overnight with membranes at 4°C. The blot was incubated for one hour at room temperature, after already being washed with an anti-mouse or anti-rabbit IgG accompanied with horseradish peroxidase (Millipore, 1:5000 dilutions in PBST containing 5% non-fat milk). Finally, the blots were visualized using a chemiluminescence detection kit (ECL; Millipore), and the densitometric scanning has quantified the optical density of each band.

Histopathological Examination

Liver morphology of Balb/c mice either normal or DEN-treated ones was examined by histopathology. Liver sections (5 micron-thickness) were fixed and stained with regular Hematoxylin-eosin stain (H&E), and then examined under light microscope and lesions were photographed with Olympus digital camera installed on the Olympus microscope with 1/2 X photo-adaptor, using 100X objective lens.

Enzyme-linked immunosorbent assay (ELISA)

Tumor necrosis factor (TNF)-α and interleukin (IL)-1β in the mouse serum were measured according to the manufacturer’s specifications (R&D Systems, Minneapolis, MN, USA). The OD value at 450 nm was determined by a Power Wave microplate reader Bio-Tek, USA.

Flow Cytometer Analysis

This analysis was performed using Annexin V FITC and Propidium Iodide (PI), in order to measure the rates of apoptosis in blood cells. Annexin V-stained flipped phospholipids in lymphocytes cell membranes and Propidium Iodide (PI)-stained that damaged DNA were used in determination of early and late apoptosis, respectively, after treatment compared to control. Annexin V and PI were purchased from (Thermo Fisher Scientific). The apoptotic analysis was dedicated to differentiate between early and late apoptotic cells, as well as necrotic cells. The Apoptosis of the treated and untreated cells was analyzed by flow cytometer instrument (Attune® Acoustic Focusing Cytometer, Life Technologies) and analyzed by its software.

Results

Mortality and occurrence of liver, lung and heart cancer or injury liver

We found high percentage of mortality (dead animals) in treated animals with DEN in either 4 or 8-weeks' treatment. In addition we found occurrence of lung and heart cancer in treated animals with DEN parallel to liver cancer (Table 2).

Histopathological Examination

Histopathological (Figures 1-2) revealed dose-dependent degenerative progression in hepatocyte structure, since administration of DEN in a dose of 50 mg/Kg once weekly i.p for 4 weeks showed fatty degeneration and scattered inflammatory cell infiltrate. While, the higher dose 75 mg/Kg for the same period of time showed steatohepatitis with fatty degeneration and scattered inflammatory cell infiltrate. Higher dose of DEN (75 mg/Kg) i.p once weekly for 8 weeks showed HCC that comprises sheets and nests of large tumor cells with pleomorphic nuclei, prominent nucleoli, abundant granular eosinophilic cytoplasm, scatter Mallory bodies (dense eosinophilic cytoplasmic structure) and distinct cell borders.
DEN stimulates expression of IQGAP1, TRAIL decoy receptors, NFKB and IL-8

The genetic expression of IQGAP1, as well as the anti-apoptotic genes TRAIL decoy receptors DcR1 and DcR2, NFKB and IL-8 and its receptors CXCR1 and CXCR2 was significantly higher (P < 0.001) in DEN-induced cancer groups, as compared to their corresponding normal control ones (Figures 3a-d and 4a). Furthermore, the expression of these genes was increased by increasing the DEN dose.

Table 1. Primer Sequences of Used Mouse Genes

| GenBank Accession | Mouse Gene | Forward Primer | Reverse Primer |
|-------------------|------------|----------------|---------------|
| NM_011399.2       | 1L-8       | CTAGGCATCTTCGTCCGTCC | AGCCCATAGTGGAGTGGGAT |
| NM_009910.3       | Cxcr3B     | TCAGCCAACTACGATCAGCG | CCTCTGGAGACCAGCAGAC |
| NM_019746.4       | DcR1       | GCTGAAGAGACAATGAAC | ACGATCACAAGGGAGGAAG |
| AB020317.1        | P53        | GCCCATGCTACAGAGGAGTC | TGAGTGGAATCTGGGATTGTG |
| NM_020275.4       | Tnfrf10b   | ACTGTCTGCCTCTGTCCTC | GACTGTAGACACACTGCTCC |
| NM_178241.4       | Cxcr1      | CAGCTGGTCAGGAAAGCAGC | CCAAGAAGGGGCGGTCAAT |
| NM_009909.3       | Cxcr2      | ATGGCTTGTCAGCAAGGAG | CTTCTACAGCAACTCCCACC |
| NM_003844.3       | DR4        | ATGAACTCTAGGTTTTCTTGGC | TTGCGTGCCCTTCCTCAAG |
| Mus musculus:DCR2 |            | AGCTGTGGTTGTGGTTGG | GGCTCAAGTTACTGACTG |
| NM_016721.2       | Igfap1     | TCTCAAGGTCTCTGCTGGGCT | GAGGCGGGTACTGGCTTAC |
| NM_027711.1       | Igfap2     | CTCAAGGTCTCATGGGCTTAC | GAGGCGGGTACTGGCTTAC |

Figure 1. Histopathological Examination. A, Normal looking of controlled mouse hepatocytes (H&E stain, 100x); B, Steatohepatitis of mouse hepatocytes treated with DEN 50 mg/kg once weekly i.p for 4 weeks showing fatty degeneration and scattered inflammatory cell infiltrate (H&E stain, 200x); C, Hepatocytes from another rat treated with DEN 75 mg/kg once weekly i.p for 4 weeks show steatohepatitis, fatty degeneration and scattered inflammatory cell infiltrate (H&E stain, 200x); D, Liver of mice treated with DEN 75 mg/kg i.p once weekly for 8 weeks shows HCC that comprises sheets and nests of large tumor cells with pleomorphic nuclei, prominent nucleoli, abundant granular eosinophilic cytoplasm, scattered Mallory bodies (dense eosinophilic cytoplasmic structures) and distinct cell borders (H&E stain, 200x).
DEN inhibits expression of IQGAP2, TRAIL death receptors, p53 and PDCD5

The genetic expression of IQGAP2, as well as the pro-apoptotic genes TRAIL death receptors DR4 and DR5, p53 and PDCD5 was significantly lower (P< 0.001) in all animal treated with DEN as compared to their corresponding normal control groups and their expressions were further decreased by increasing the DEN dose as shown in Figures 3a- d and 4a).

Protein analysis

Protein level of IQGAP1 showed a highly significant increase in DEN-treated group (4 weeks) as compared to normal controls while p53 protein level was significantly decreased in DEN-treated group as compared to the control (Figure 4b-c).

Enzyme-linked immunosorbent results

Both of tumor necrosis factor (TNF)-α and interleukin (IL)-1β were highly significant increase in both treated animals with DEN if compared with negative control.

Detection of early and late apoptosis by Flow Cytometer

All treated animals with DEN either 4 or 8 –weeks showed high incidence of early and late apoptosis if compared with negative control. In 8- weeks treated group, there were significant increases in necrotic cells if compared to 4- weeks treated group (Figure 5).

Discussion

IQGAP family of proteins is known to possess various protein-protein interaction domains. These members control certain cellular processes, including cell adhesion and migration, cytokinesis and many extracellular signals. Three IQGAPs occur in human beings: IQGAP1 and IQGAP2. IQGAP1 is this family’s best characterized protein and is known to be over-expressed in HCC (Zoheir et al., 2015). This IQGAP1 has a domain of calponin homology, a domain of polyproline, and a RasGAP-related domain (GRD), as well as a sequence of RasGAP carboxy-terminal that is common to members of the IQGAP family (Brown and Sacks, 2006). IQGAP1 assists in both ERK-and β-catenin-dependent signaling. IQGAP1 binds B-Raf, MEK and ERK to facilitate their sequential activation and propagation of the MAPK cascade. Constitutive activation of the MAPK pathway is a common oncogenic trigger in several cancers (White et al., 2009).

The process of programmed cell death is initiated in cells in response to either extracellular cues via stimulation of the death receptors (DRs), often referred to as the extrinsic pathway, or to a variety of intracellular stress conditions that involve mitochondria and are known as the intrinsic pathway. A distinct mode of programmed apoptosis is critical for tissue homeostasis and its deregulation is frequent in malignancy. The extrinsic pathway starts with binding of death ligands to
Figure 3. DEN Regulates Expression of IQGAPs, DcR1 and DcR2, DR4, DR5 and NF-κB Genes in Liver Tissues.

a, mRNA amounts of IQGAP1 and 2 were quantified using RT-PCR and normalized to GAPDH housekeeping gene as in Methods; b, mRNA amounts of DcR1 and DcR2 were quantified using RT-PCR and normalized to GAPDH housekeeping gene as in Methods. Statistical analysis was performed using ANOVA followed by t-test. Each value indicates the mean ± S.E.M of ten animals. p < 0.05 was considered significant (*) compared to the corresponding normal control (0 mg/kg) group.

Figure 4. a, DEN suppresses expression of IL-8, CXCR1 and CXCR2 genes in liver tissues; b and c, DEN upregulates IQGAP1 and downregulates p53 expression in DEN-treated liver cells. Balb/c mice were treated with noted concentrations of DEN 4 weeks. Liver tissues were homogenized and total protein was prepared as described in Methods. IQGAP1 and p53 protein levels were determined by Western blot analysis (upper panel). Representative densitometric quantifications of IQGAP1 and p53 protein levels are shown in the lower two panels. Values are presented as means ± standard error of the mean. *p<0.05, compared with control (NC).
Figure 5. Flow Cytometry Analysis Using Annexin V FITC and Propidium Iodide (PI) for Apoptosis Measurements. Blood cells were treated with DEN. Early, late apoptosis, and necrosis were detected in the stained cells with Annexin V and PI.

According to our results, the expression level of IQGAP1 was significantly upregulated in HCC tissues, compared to normal liver tissue. This corroborates previous findings by Chen et al., (2010), who reported that IQGAP1 is overexpressed in HCC. Our data also show that NF-kB and IL-8 and its receptors were upregulated in HCC tissues, as compared to normal ones. These data confirm previous findings showing that NF-kB (He and Karin, 2011) and IL-8 (Ren et al., 2003) are upregulated in HCC. Knowing that IQGAP1 triggers activation of NF-kB (Tseng et al., 2014), our data reveals that IQGAP1 stimulates NF-kB signaling, which in turn, augments expression of its downstream effectors IL-8 and its receptors CXCR1 and CXCR2. Notably, to the best of our knowledge, we are the first group to report overexpression of IL-8 receptors in DEN-induced HCC and the first group to report overexpression of CXCR2 in HCC, in general. Moreover, we report, for the first time, that DEN activates such an IQGAP1/ NF-kB/IL-8 pathway in HCC in vivo.

The pro-apoptotic TRAIL death receptors were markedly downregulated in HCC cells, as compared to normal liver cells. In addition, DEN induced upregulation of TRAIL decoy receptors. This suggests that this regulation of TRAIL receptors facilitated DEN-induced mutagenicity and carcinogenicity. Since our data showed that DEN stimulated NF-kB and IL-8 expression, and NF-kB and IL-8 are known to suppress TRAIL-induced apoptosis, we may conclude that DEN fostered NF-kB expression, which in turn upregulated IL-8 expression and then both NF-kB and IL-8 suppressed TRAIL death receptors expression. In addition, TRAIL death receptors are known to be transcriptionally upregulated by p53 (Willms et al., 2019). Since p53 has been shown to inhibit NF-kB activity (Shao et al., 2000) and our data reveal that DEN significantly decreases p53 mRNA and protein levels, this could allow for upregulation NF-kB,
which in turn could stimulate IL-8 signalling and inhibit TRAIL death receptors expression. Furthermore, since NF-kB is known to upregulate TRAIL decoy receptors (Bernard et al., 2001), we may conclude that DEN-induced upregulation of NF-kB may have augmented expression of TRAIL decoy receptors.

Our data showed that PDCD5 was downregulated in HCC tissues, relative to normal liver tissue. This agrees with findings by Fu et al., (2013) who reported that expression of PDCD5 was lower in HCC tissue compared to normal tissue. Knowing that PDCD5 is an important upstream regulator of P53 which upregulates p53 (Fu et al., 2013), we may conclude that DEN-induced PDCD5 downregulation in HCC resulted in diminished expression of p53, which then allowed for stimulation of the IQGAP1/ NF-kB /IL-8 pathway.

In conclusion, our DEN mouse model for liver cancer clearly showed that that there is a crosstalk between these two pathways, namely IQGAP1/NF-kB/IL-8 and PDCD5/ p53/TRAIL/caspase-8/caspase-3 pathways. Occurrence of apoptosis and cell death and necrosis were confirmed via other parameters such as flow cytometer and tumor necrosis factor (TNF)-α and interleukin (IL)-1β. In addition, DEN made lung and heart cancer parallel to liver cancer which will be the new topic for further manuscript.

**Author Contribution Statement**

All authors participated in the design, interpretation of the studies and analysis of the data and review of the manuscript; KZ designed this study. KM, MA, AD, and KZ participated in the conduct of the study. MA, AD, KM and K.Z conducted the experiments, KZ supplied critical reagents, KZ wrote the manuscript.

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**Ethics Approval and Consent to Participate**

The experimental protocol used in the study was approved (NRCE-CBD1032019) by the Animal Care and Use Committee of National Research Centre, Egypt.

**Availability of data and material**

Data is available for all.

**Competing interests**

There is no competing interest.

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