Inhibition of Polyprotein Processing and RNA Replication of Human Rhinovirus by Pyrrolidine Dithiocarbamate Involves Metal Ions

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Pyrrolidine dithiocarbamate (PDTC) is an antiviral compound that was shown to inhibit the replication of human rhinoviruses (HRVs), poliovirus, and influenza virus. To elucidate the mechanism of PDTC, the effects on the individual steps of the infection cycle of HRV were investigated. PDTC did not interfere with receptor binding or internalization by receptor-mediated endocytosis of HRV2 particles into HeLa cells. But we demonstrate that the processing of the viral polyprotein was prevented by PDTC treatment in HeLa cells infected with HRV2. Furthermore, PDTC inhibited the replication of the viral RNA, even when added four hours post infection. As PDTC is described as a metal ion binding agent, we investigated the effect of other metal chelators on the multiplication of HRV2. We show that EDTA, o-phenanthroline, and bathocuproine disulfonic acid do not exhibit any antiviral properties. Surprisingly, these substances, coadministered with PDTC, abolished the antiviral effect of PDTC, suggesting that metal ions play a pivotal role in the inhibition of virus multiplication. These results suggest that PDTC inhibits the activity of the viral proteases in a metal ion dependent way.

Human rhinoviruses (HRVs) are the most frequent cause of the common cold and are implicated in more than 50% of upper respiratory tract infections (45). Although not life threatening, infection with HRV can prepare the ground for more serious diseases, such as acute exacerbation of asthma (20, 30) or otitis media (5). As there are more than 100 HRV serotypes, vaccine development is unfeasible. Apart from symptomatic medication no causative treatment for HRV infections is currently available. Therefore, analysis of functions of new antiviral substances is of great interest and might also demonstrate that the processing of the viral polyprotein was prevented by PDTC treatment in HeLa cells infected with HRV2. Furthermore, PDTC inhibited the replication of the viral RNA, even when added four hours post infection. As PDTC is described as a metal ion binding agent, we investigated the effect of other metal chelators on the multiplication of HRV2. We show that EDTA, o-phenanthroline, and bathocuproine disulfonic acid do not exhibit any antiviral properties. Surprisingly, these substances, coadministered with PDTC, abolished the antiviral effect of PDTC, suggesting that metal ions play a pivotal role in the inhibition of virus multiplication. These results suggest that PDTC inhibits the activity of the viral proteases in a metal ion dependent way.

Ribosome entry site (RES), which is located in the 5′ untranslated region of the HRV genome. The polyprotein is processed into the mature viral proteins through a sequence of cleavages performed by two virus-encoded proteases. The 2A proteinase (2Apro) cleaves between the C terminus of VP1 and its own N terminus to separate the capsid protein region from the nonstructural protein precursor (26, 47). Promising clinical trials have been performed with the capsid binding substance pleconaril (34) and 2Apro inhibitor rupintrivir (AG7088) (31).

Viral replication via a minus-strand intermediate and subsequent synthesis of new positive-strand RNA is mediated by the virus-encoded RNA-dependent RNA polymerase 3Dpol.

Finally, RNA and capsid proteins are assembled into mature infectious viral particles, which are then released by cell destruction. During late stages of virus infection morphological changes of cells can be observed, known as cytopathic effects.

Over the last two decades several steps in the life cycle of HRV have been targeted for antiviral therapy (39), such as viral attachment, uncoating, polyprotein processing, and RNA polymerization. So far, only limited success has been achieved by the use of recombinant soluble receptor fragments to block virus binding (26, 47). Promising clinical trials have been performed with the capsid binding substance pleconaril (34) and the 3Dpol inhibitor rupintrivir (AG7088) (31).

Recently, we described pyrrolidine dithiocarbamate (PDTC) as a potent inhibitor of HRV multiplication. PDTC is effective against all tested HRV serotypes in several cell lines analyzed without being toxic (11). In HeLa cells, the IC 50 against HRV...
14 is 60 μM, toxic effects can be observed at concentrations higher than 1 mM. Surprisingly, the antiviral activity of PDTC is not restricted to members of the Picornaviridae such as HRV or poliovirus, since PDTC was also shown to inhibit multiplication of influenza virus, a member of the Orthomyxoviridae (48; A. Grassauer, personal communication). Therefore, the antiviral activity of this compound seems to involve more general processes unidentified so far.

PDTC is a low-molecular-weight thiol compound that can exert numerous effects in biological systems. It is a widely used inhibitor of the activation of nuclear factor kappa B (NF-κB) (37, 38), although there are conflicting reports how this inhibition is accomplished. On the one hand it has been attributed to the antioxidative property of PDTC (2), on the other hand to the inhibition of the ubiquitin ligase activity, which is a prerequisite for NF-κB activation (16). Furthermore, modulation of the cellular availability of heavy metal ions is proposed as another mechanism of the inhibition of NF-κB activation by PDTC (2, 38). PDTC binds various metal ions, leading to the formation of lipophilic dithiocarbamate-metal complexes, which transport metal ions such as zinc and copper from the extracellular medium into the cell (9, 21, 43).

To gain insight into the mechanism by which PDTC blocks the replication of HRV, we examined the effects on the various steps of the viral life cycle in detail. Receptor binding and internalization of HRV2 are not affected by PDTC. However, we demonstrate that PDTC blocks processing of the viral polyprotein. Addition of PDTC, even at 4 h postinfection, results in a total inhibition of replication of both positive- and negative-strand RNA.

As PDTC is a metal binding substance and several viral processes were described to be influenced by various metals, we investigated if the metal ion binding function of PDTC is the mechanistic basis of the antiviral function. Interestingly, the antiviral action of PDTC is abolished by the addition of metal chelating agents, such as EDTA, α-phenanthroline, or bathocuproine disulfonic acid (BCS) and can be restored by the concomitant addition of zinc and copper ions. These results suggest that metal ions play an important role for the antiviral property of PDTC.

**MATERIALS AND METHODS**

**Cells, media, and reagents.** HeLa cells (strain Ohio; European Collection of Cell Cultures, Salisbury, United Kingdom) were cultured in RPMI 1640 (Invitrogen) supplemented with 10% heat-inactivated fetal calf serum (Invitrogen), 2 mM l-glutamine (Dipro, Austria), 100 U/ml penicillin (Dipro), and 100 μg/ml streptomycin (Dipro).

Chemicals were obtained from Sigma. PDTC was purchased from Alexis Biochemicals.

**Virus preparation.** HRV2 was obtained from the American Type Culture Collection and routinely grown in suspension cultures of HeLa cells (strain Ohio; Flow Laboratories, McLean; Virginia) as described previously (40). Virus titers in 50% tissue culture infectious doses (TCID₅₀)/ml were determined according to Reed and Muench (33).

**Infection of cells.** HeLa cells seeded into 6-well tissue culture dishes were cultured in RPMI medium supplemented with 2% heat-inactivated fetal bovine serum, 2 mM l-glutamine, 100 U/ml penicillin, and 100 μg/ml streptomycin (infection medium). PDTC was added simultaneously with HRV. Routinely, 125 μM PDTC was used, which is nontoxic to HeLa cells (11). Cells were incubated at 37°C and 5% CO₂ in a humidified incubator. The input virus was removed 2 h postinfection, and cells were washed once with an acidic HEPES buffer (10 mM HEPES, 140 mM KCl; pH 5.3) and twice with phosphate-buffered saline (PBS). For further incubation, infection medium and PDTC were added when indicated.

**Immunofluorescence.** Immunofluorescence was essentially performed as described elsewhere (3). Briefly, cells were grown on sterile coverslips in six-well tissue culture dishes. Fixation was done with 4% paraformaldehyde–PBS at room temperature (RT) for 10 min. Cells were permeabilized by incubation in 0.5% Triton X-100 in PBS for 5 min. Blocking was done at 4°C using 1% goat serum–PBS. As primary antibody the mouse monoclonal antibody 8F5 (1:200 in PBS–1% bovine serum albumin) against the virus capsid protein VP2 (1:200) was employed (29, 40). As secondary antibody, Alexa680-labeled goat anti-mouse immunoglobulin G conjugates were used at a dilution of 1:800 (Molecular Probes). Finally, slides were mounted with DAKO fluorescent mounting medium and examined with a Leica TCS-NT confocal microscope using fluorescein iso-thiocyanate settings.

**Measurement of IRES-dependent translation.** HeLa cells were infected with HRV2 using the multiplicity of infection indicated in the figure legends. The culture medium was removed 6 h postinfection, the cells were pretreated with 125 μM PDTC or 125 μM cycloheximide for 15 min and washed twice with PBS. The cells were labeled in 1 ml per well of methionine- and cysteine-free RPMI medium supplemented with 300 μCi of [35S]methionine/cysteine (radiochemical purity by high-pressure liquid chromatography: 70% [35S]methionine, 25% [35S]cysteine; Hartmann Analytic, Braunschweig, Germany) for 1 h. The labeling was performed in the presence of PDTC or cycloheximide. Subsequently, proteins were precipitated with trichloroacetic acid, and the incorporated radioactivity was measured in a scintillation counter (Packard).

**Pulse-labeling and immunoprecipitation.** HeLa cells were infected with HRV2. The infection medium was replaced by 1 ml per well of methionine- and cysteine-free RPMI medium supplemented with 600 μCi of [35S]methionine/cysteine (radiochemical purity by high-pressure liquid chromatography: 70% [35S]methionine, 25% [35S]cysteine; Hartmann Analytic, Braunschweig, Germany) at 6 h or 6.5 h postinfection During a labeling period of 1 h 125 μM PDTC, or 200 μM ZAV.fmk (benzoxolonylcarbonyl-Val-Ala-Asp-fluoromethyl ketone) were added. Protein extracts were prepared by addition of reducing sample buffer (20% glycerol, 2% mercaptoethanol, 8% sodium dodecyl sulfate [SDS], 0.05% sodium decyl sulfate blue and examined by SDS-polyacrylamide gel electrophoresis (PAGE). Alternatively, the cells were incubated in infection medium supplemented with PDTC.

For immunoprecipitation the cells were washed twice with PBS, collected by scraping in 1 ml of ice-cold PBS and centrifugation at 300 × g for 4 min. Cell pellets were frozen in liquid N₂ and stored at −20°C. They were resuspended by vortexing in 500 μl of lysis buffer (1% [w/v] Triton X-100, 50 mM Tris-Cl, 300 mM NaCl, 5 mM EDTA, 0.02% Na-azide; pH 7.4) supplemented with 10 mM iodoacetamide, 1 mM phenylmethylsulfonyl fluoride, and 2 μg/ml leupeptin. The samples were incubated on ice for 30 min and centrifuged at high speed in an Eppendorf centrifuge at 4°C for 15 min. Meanwhile, 1 μg of the monoclonal antibody 8F5 specific for VP2 was coupled to 30 μl 50% pNP protein A-Sepharose bead slurry (Amersham Bioscience) in 0.5 ml ice-cold PBS and 0.01% Triton X-100. The beads were washed three times with lysis buffer; 500 μl of cell lysates were added, and the antibody-bound beads were combined and tumbled at 4°C for 2 h. The protein-bead conjugates were washed twice with ice-cold PBS. Proteins were resuspended in sample buffer, separated by SDS-PAGE and visualized by exposure of the dried gel to BioMax MR x-ray film (Kodak).

**Northern blotting.** Total RNA was extracted from infected HeLa cells using RNeasy mini kit (QIAGEN). 10 μg of each sample were fractionated on a 1% formaldehyde denaturing gel at 60 V, and the RNA quality was checked by ethidium bromide staining to visualize the 18S and 28S rRNAs. RNA was purified by high-pressure liquid chromatography: 70% [35S]methionine, 25% [35S]cysteine; Hartmann Analytic, Braunschweig, Germany) using 1. Membranes were incubated in prehybridization solution consisting of 25 ml 20x SSC (350 mM NaCl plus 0.15 M sodium citrate) buffer overnight and UV-cross-linked (120 kJoule) in a Stratalinker (Stratagene). For synthesis of strand-specific RNA probes, primer extension was performed in the presence of [γ-32P]dCTP (purity >98%; Hartmann Analytic, Braunschweig, Germany) using the cDNA of HRV2 as a template (41). The primers for the positive and negative strands were 5′-TGCCGCTTGGAGATATAGCGT-3′ (located in the 2C coding region) and 5′-CAGCTAGGGTGTAATAGTGCT-3′ (located in the 3C coding region), respectively. The primer extension reaction was set up as follows: 0.2 mM dATP, 0.2 mM dTTP, 0.2 mM dGTP, 2 μM dCTP, 54 μCi of [γ-32P]dCTP, 25 pmol primer, 2.5 U Taq polymerase (Finnzymes, Espoo, Finland), 5 ng of HRV2 template in reaction buffer in a total volume of 50 μl. Membranes were incubated in prehybridization solution consisting of 25 ml 20x SSPE buffer (2 M NaCl, 0.25 M NaH₂PO₄, 0.02 M EDTA; pH 7.4), 50 ml 100% formamide, 10 ml blocking reagent (Roche), 1 ml 10% SDS, and H₂O to 100 ml at 42°C for 2 h. Labeled RNA probes were denatured at 95°C for 5 min and added to the prehybridizing solution. Mem-
PDTC does not interfere with internalization of HRV2. 

HeLa cells grown on coverslips were infected with HRV2 (multiplicity of infection, 50) in the absence or presence of 125 μM PDTC. Cells were fixed using 4% paraformaldehyde 20 min, 60 min, and 8 h postinfection. The virus capsid protein VP2 was localized by the monoclonal antibody 8F5 and visualized by immunofluorescence.

RESULTS

PDTC does not alter the infectivity of HRV particles. Over the past two decades, antirhinoviral agents inhibiting viral attachment and/or uncoating were developed, e.g., WIN compounds and pleconaril (4). To find out whether PDTC interacts with the viral capsid, HRV2 particles were incubated with PDTC at room temperature for 1 h. Afterwards, HeLa cells were incubated with PDTC-treated or untreated virus at 4°C for 1 h. Under these conditions, receptor binding but no further internalization can take place. Unbound virus was removed by extensive washing and infection was continued by cultivating cells in infection medium without PDTC at 37°C. The virus titer of the supernatant was determined by titration using TCID₅₀ assays 24 h postinfection.

Infection of HeLa cells with PDTC-treated virus resulted in a titer of 1.8 × 10⁶. A similar titer was obtained in control infections with untreated virus (Fig. 1). In contrast, continuous presence of PDTC during infection led to a significant reduction of the virus titer of three orders of magnitude (Fig. 1) (11). Thus, PDTC inhibits the multiplication of HRV2 when present during the infection, but it does not directly interact with or inactivate HRV2 particles.

Receptor binding and internalization of HRV2 is not affected by PDTC. HRV enters the cell by receptor-mediated endocytosis. Minor group serotypes like HRV2 bind to members of the low-density lipoprotein receptor family (18). To investigate the effect of PDTC on receptor binding and internalization of HRV2, the localization of the virus particles during infection was monitored by immunofluorescence staining of the virus capsid protein VP2.

An infection period of 20 min resulted in a uniform distribution of the HRV2 particles in the cytoplasm of the HeLa cells (Fig. 2a); 60 min postinfection the virus accumulated in the perinuclear region, which is the characteristic localization of late endosomes. HeLa cells infected with HRV undergo morphological changes, like cell rounding and detachment from the substrate, described as cytopathic effect. As a complete life cycle of HRV2 in HeLa cells lasts approximately 10 h, the cytopathic effect became visible 8 h postinfection, and the cytoplasm of the infected cells were filled with progeny viral particles (Fig. 2c).

In the presence of PDTC the amount and distribution of HRV2 particles was similar to that of nontreated cells at 20 and 60 min postinfection (Fig. 2). Thus, the initial steps of virus infection such as receptor binding and internalization are not affected by PDTC.

However, treatment with PDTC completely inhibited the multiplication of viral particles at 8 h postinfection, and later on a loss of intracellular viral particles was observed (Fig. 2 and data not shown). In addition, PDTC prevented the formation of a visible cytopathic effect.

PDTC has no major effect on IRES-dependent translation. During rhinoviral infection the cellular cap-dependent translation is inhibited by the host cell shut-off. This is achieved by cleavage of the cellular translation factors eIF4GI and eIF4GII by the viral protease 2Apro (10, 14). However, translation of the HRV RNA genome is unaffected, as it is initiated at an internal ribosomal entry site (IRES) via a cap-independent mechanism. In the presence of PDTC no significant synthesis of viral proteins can be detected (see Fig. 2A in reference 11). Among other reasons, this can be due to decreased viral translation or impaired polyprotein processing.

To investigate the effect of PDTC on IRES-dependent translation the level of protein synthesis was determined after onset of host cell shut-off in cells infected with HRV2 by incorporation of [³⁵S]methionine/cysteine for 1 h followed by trichloroacetic acid precipitation of the proteins and scintillation counting. During the labeling period cells were treated with PDTC or cycloheximide, a strong inhibitor of both cap- and IRES-dependent translation. PDTC slightly reduced the level of IRES-dependent translation (about 80% compared to untreated cells), whereas cycloheximide significantly decreased
The level of translation to about 20% (Fig. 3). This weak effect of PDTC on IRES-dependent translation is unlikely to be responsible for the absence of newly synthesized capsid proteins.

**PDTC interferes with polyprotein processing of HRV2.** Upon infection, the HRV genome is translated into a large polyprotein precursor, which is subsequently cleaved by a sequence of proteolytic cleavages. To investigate the effect of PDTC on the processing of the viral polyprotein, we analyzed the pattern of virus protein precursors in the presence and absence of PDTC by pulse-labeling. As a positive control we employed the methylated form of the caspase inhibitor zVAD.fmk, which was recently described to inhibit HRV 2Apro activity of several viral enzymes was described to be influenced by metal ions, we employed metal-chelating substances. In this context it has to be noted that several biological effects of agents.

In untreated cells an increase in both positive- and negative-strand RNA levels was detected starting 4 h postinfection, which resulted in a peak RNA level at 8 h postinfection (Fig. 5). At 13 h postinfection the level of viral RNA decreased again due to the cytopathic effect. In contrast, addition of PDTC at the onset of viral RNA synthesis 4 h postinfection gave rise to an immediate and strong inhibition of both positive and negative strand RNA replication (Fig. 5). Viral RNA synthesis is also reduced to the same extent when PDTC is administered at the time of infection (data not shown).

Thus, PDTC strongly inhibits viral RNA synthesis of both positive and negative strands. A repression of replication of virus RNA by PDTC was also observed for the major group virus HRV14 (data not shown).

**Antiviral effect of PDTC is abolished by metal ion chelating agents.** Previously, Erl et al. showed that PDTC can increase the intracellular concentration of various metal ions (9). As the activity of several viral enzymes was described to be influenced by metal ions, we employed metal-chelating substances. In this context it has to be noted that several biological effects of PDTC, such as inhibition of the activation of NF-κB or induction of apoptosis in endothelial cells, are abolished in the presence of metal ion chelators (21, 22).

To investigate whether the antiviral effect of PDTC was dependent on its ability to bind metal ions, other metal ion chelators such as o-phenanthroline, BCS, or free-base EDTA were added during virus infection. The amount of progeny virus in the supernatant was determined by TCID_{50} assays 24 h containing intermediate precursor proteins were immunoprecipitated with the monoclonal antibody BFS (40), and analyzed by SDS-PAGE and autoradiography. To clearly identify VP2, bands were compared to radiolabeled HRV 2 virus separated simultaneously (lane *HRV2*) (28).

In untreated cells the processing of the P1 precursor protein was clearly visible as a reduction of the amount of the corresponding polyprotein band. At the same time a smaller band comprising VP2 appeared (Fig. 4B). In the presence of PDTC, the processing of the P1 protein was blocked and no cleavage products could be detected.

Furthermore, we were interested whether the inhibitory effect of PDTC on the processing of the polyprotein does depend on de novo protein synthesis. To this end, proteins of infected cells were labeled for 15 min starting at 6 h 45 min postinfection. Afterwards, the cells were treated with PDTC and/or cycloheximide for 2 h. Again, virus proteins were analyzed by SDS-PAGE and autoradiography.

In untreated cells the large precursor molecules were processed into the various viral proteins (Fig. 4C). Cycloheximide did not interfere with the viral polyprotein processing. However, P1 protein processing was impaired in the presence of PDTC, resulting in P1 accumulation. When PDTC and cycloheximide were coadministered, the pattern of the virus proteins was similar to that of PDTC-treated cells alone. This indicates that the inhibitory effect of PDTC on polyprotein processing does not depend on de novo protein synthesis.
postinfection. The multiplication of HRV2 was not affected by these metal chelators at the concentrations used, which is in agreement with published data, where 1,000-fold higher concentrations of EDTA were shown to interfere with receptor binding (data not shown) (25). However, coadministration of metal chelators and PDTC abolished the antiviral effect of PDTC. As shown in Fig. 6 the virus titer of HRV2 was reduced by PDTC alone about three orders of magnitude compared to controls. When BCS was added simultaneously the antiviral effect was abolished in a dose-dependent manner, leading to normal virus replication as in untreated cells at 1,000 μM BCS (Fig. 6A). Similar effects were observed after coadministration of EDTA or -phenanthroline (Fig. 6B and C). EDTA was much more potent in terms of inhibiting the effect of PDTC, as 10 μM EDTA was already sufficient compared to 62 μM -phenanthroline. Interestingly, both the cell-permeable agent -phenanthroline and the non-cell-permeable substances EDTA and BCS were effective.

Copper and zinc ions are involved in the antiviral effect of PDTC. Several divalent metal ions, including copper and zinc, have been shown to contribute to some of the biological effects of PDTC, such as the inhibition of NF-κB and induction of cell apoptosis (6, 9, 21, 22). We analyzed whether zinc, iron, or copper ions could restore the antiviral activity of PDTC when it is suppressed by EDTA (as shown in Fig. 6). Addition of at least 2.5 μM Zn2+ ions or 25 μM Cu2+ ions reestablished the antiviral activity of PDTC in the presence of EDTA (Fig. 7A and C). Iron ions failed to show an effect (Fig. 7B). None of these metal ions alone showed any antiviral activity in the concentrations used (Fig. 7) and no cytotoxicity was observed (data not shown). These results suggest that Zn2+ or Cu2+ ions are an important factor for the antiviral effect of PDTC.

**DISCUSSION**

Currently, there is no cure available for the common cold apart from symptomatic treatment. PDTC was shown to be a potent inhibitor of the multiplication of HRV and poliovirus, both members of the Picornaviridae (11). Furthermore, PDTC turned out to prevent the multiplication of influenza virus, belonging to the Orthomyxoviridae (48). This indicates that the
The antiviral effect of PDTC is not restricted to one distinct virus family, and thus its mode of action might rely on more general mechanisms. The precise molecular basis of the inhibitory function has not yet been determined.

To elucidate the action of PDTC on HRV multiplication in more detail, we investigated the effect of PDTC on single steps of the infection cycle of HRV2. We demonstrated that PDTC did not affect the early steps of the viral infection cycle, such as receptor binding, virus entry, and internalization. Thus, PDTC does not interact with the capsid of HRV2, as preexposure of the virus to PDTC did not alter the infectivity of HRV particles (Fig. 1). Moreover, PDTC does not interfere with virus entry, since binding to the cellular receptor and receptor-mediated endocytosis were unaffected by PDTC (Fig. 2). Although PDTC slightly decreased IRES-dependent translation, this effect is too weak to be exclusively responsible for the antiviral property of PDTC (Fig. 3).

However, our data indicate that PDTC impedes the processing of the HRV polyprotein (Fig. 4). As PDTC treatment abolished the appearance of processed VP2 an inhibition of the viral proteases 2Apro and/or 3Cpro is proposed (Fig. 4B). As a consequence of the impaired polyprotein processing, the repressed replication of both positive- and negative-strand RNA in the presence of PDTC might be due to alterations in the processing and production of the nonstructural proteins (Fig. 5).

In account of their important roles in the processing of the
viral polyprotein, blocking of the viral proteases 2A\textsuperscript{pro} and 3C\textsuperscript{pro} is regarded as a promising approach to antiviral therapy (27, 35, 45). Recently, we demonstrated that inhibition of the viral protease 2A\textsuperscript{pro} with the methylated form of the caspase inhibitor zVAD.fmk resulted in a significant decrease of multiplication of HRV (8). The most promising HRV protease inhibitor currently available is the 3C\textsuperscript{pro} inhibitor pruninivir, which shows potent activity for a broad spectrum of rhinovirus serotypes (31).

Previously, we have described that PDTC does not inhibit the activity of the HRV2 2A\textsuperscript{pro} in vitro, as was shown by using purified protease and cleavage of synthetic peptide substrates (11). Furthermore, PDTC failed to inhibit 2A\textsuperscript{pro}-mediated cleavage of eIF4GI and cytokeratin 8 in cell extracts. No inhibition of eIF4GI cleavage was found in rabbit reticulocyte lysates during translation of mRNAs encoding HRV2 2A\textsuperscript{pro} (data not shown). Nevertheless, in a cellular context PDTC might change the cytoplasmic environment, thereby interfering with 2A\textsuperscript{pro} and/or 3C\textsuperscript{pro} function.

Dithiocarbamates can chelate various metal ions, leading to the formation of lipophilic dithiocarbamate-metal complexes (43). Several studies have reported that dithiocarbamates can promote cell entry of metals, e.g., diethyldithiocarbamate and PDTC increased the intracellular copper levels in thymocytes (9). Kim et al. demonstrated that PDTC gave rise to uptake of zinc ions (22). These data suggest that PDTC might alter the tightly regulated balance of ions in cells, resulting in alterations of enzyme functions. In some reports it was described that various biological effects of PDTC can be prevented by the addition of metal chelators (21, 22). In this manuscript we demonstrate that the antiviral effect of PDTC was neutralized by addition of free-base EDTA, o-phenanthroline, or BCS. Thus, it seems that the availability of ions is crucial for the antiviral action of PDTC (Fig. 6).

Among metal ions, zinc has been known to interfere with HRV replication for many years. As a mechanism, the inhibition of the posttranslational cleavage of the precursor protein was suggested (23, 24). Cordingley et al. described that 3C\textsuperscript{pro} is inhibited by zinc ions in an in vitro protease assay (7). In contrast, there is no effect of zinc ions on the 2A\textsuperscript{pro} self-processing or cleavage of eIF4GI determined by in vitro translation of HRV2 mRNA in rabbit reticulocyte lysate (13).

Comparison of the antiviral activities of various zinc salts showed that the therapeutic index is very low (12). There are conflicting results on the effectiveness of zinc gluconate or zinc acetate in the treatment of the common cold in clinical trials (42, 46). However, it cannot be excluded that PDTC has an impact on viral enzymes other than proteases. Zinc ions are described to inhibit the activity of the 3D\textsuperscript{pol} of HRV16 in vitro (19). Thus, by increasing the intracellular zinc ion concentration PDTC might directly inhibit polymerase function and replication.

In our report, we propose a model for the antiviral effect of PDTC on HRV. We demonstrated that PDTC interferes with polyprotein processing. Moreover, the level of virus RNA was decreased in PDTC-treated cells. However, our results exclude an action of PDTC on the early steps of virus infection, such as receptor binding or internalization. Since the antiviral effect of PDTC is abolished in the presence of several metal ion-chelating compounds, the antiviral activity of PDTC seems to depend on the availability of metal ions. Specifically, zinc or copper ions seem to play a crucial role, as they restore the antiviral property of PDTC in the presence of EDTA.

However, to date it is unclear if PDTC works as a carrier for the ion or if binding of an ion is necessary to allow PDTC to enter the cell. Elucidation of the antiviral mechanism of PDTC will be the subject of further research and will provide more information about the environmental requirements for the rhinovirus life cycle.

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ANTIVIRAL ACTIVITY OF PDTC 13899