Distinct reaction pathway promoted by non-divalent-metal cations in a tertiary stabilized hammerhead ribozyme

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ABSTRACT

Divalent ion sensitivity of hammerhead ribozymes is significantly reduced when the RNA structure includes appropriate tertiary stabilization. Therefore, we investigated the activity of the tertiary stabilized “RzB” hammerhead ribozyme in several nondivalent ions. Ribozyme RzB is active in spermidine and Na\(^+\) alone, although the cleavage rates are reduced by more than 1,000-fold relative to the rates observed in Mg\(^{2+}\) and in transition metal ions. The trivalent cobalt hexammine (CoHex) ion is often used as an exchange-inert analog of hydrated magnesium ion. Trans-cleavage rates exceeded 8 min\(^{-1}\) in 20 mM CoHex, which promoted cleavage through outersphere interactions. The stimulation of catalysis afforded by the tertiary structural interactions within RzB does not require Mg\(^{2+}\), unlike other extended hammerhead ribozymes. Site-specific interaction with at least one Mg\(^{2+}\) ion is suggested by CoHex competition experiments. In the presence of a constant, low concentration of Mg\(^{2+}\), low concentrations of CoHex decreased the rate by two to three orders of magnitude relative to the rate in Mg\(^{2+}\) alone. Cleavage rates increased as CoHex concentrations were raised further, but the final fraction cleaved was lower than what was observed in CoHex or Mg\(^{2+}\) alone. These observations suggest that Mg\(^{2+}\) and CoHex compete for binding and that they cause misfolded structures when they are together. The results of this study support the existence of an alternate catalytic mechanism used by nondivalent ions (especially CoHex) that is distinct from the one promoted by divalent metal ions, and they imply that divalent metals influence catalysis through a specific nonstructural role.

Keywords: catalytic mechanism; cobalt hexammine; divalent cations; hammerhead ribozymes

INTRODUCTION

Hammerhead ribozymes catalyze site-specific cleavage and ligation reactions in the replication cycles of some plant viroid satellite RNAs, and they have been found in several eukaryotic genomes. The minimal motif consists of three helices that intersect at a conserved catalytic core of 11 nucleotides (nt). In crystallographic and solution studies, these minimal hammerhead ribozymes assume a Y-shaped conformation, with stems I and II making a close approach (Pley et al. 1994; Tuschl et al. 1994; Scott et al. 1995; Penedo et al. 2004). Under standard in vitro assay conditions (10 mM Mg\(^{2+}\)), a well-behaved, minimal hammerhead ribozyme can be expected to cleave its substrate at a rate of \(~1\) min\(^{-1}\). Intracellular activity is typically modest or poor, largely due to low intracellular Mg\(^{2+}\) concentrations. Several studies during the last several years highlight the importance of peripheral sequences in natural and engineered hammerhead ribozymes for their in vitro activity under physiological Mg\(^{2+}\) conditions and for their activity inside cells (De la Pena et al. 2003; Khorova et al. 2003; Saksmerprome et al. 2004; Yen et al. 2004). These peripheral elements pull the ribozyme into a compact structure that aligns the scissile phosphate for in-line displacement and that positions nucleobases G8 and G12 for acid–base chemistry (Martick and Scott 2006), as has been suggested by cross-linking studies and analysis of the pH dependence of cleavage kinetics (Han and Burke 2005; Heckman et al. 2005; Lambert et al. 2006a,b).

Site-specific Mg\(^{2+}\) binding within minimal hammerhead ribozymes has been suggested by crystallographic analysis,
electron pair resonance studies, hydroxyl radical footprinting analysis, and thiophilic metal ion rescue studies (Scott et al. 1995; Wang et al. 1999; Hanssicker and DeRose 2000; Maderia et al. 2000; Hampel and Burke 2003; Vogt et al. 2003). A Mg\(^{2+}\)-binding site adjacent to the tandem G•A mismatches is present in all the reported X-ray structures for minimal hammerhead ribozymes (Wedekind and McKay 1998), although an active site metal ion is not evident in the native (extended) Schistosoma structure (Martick and Scott 2006). Another specific metal binding site adjacent to the phosphodiester cleavage site has been implicated from thiophilic substitution studies of the non-bridging pro-Rp oxygen of the cleavage site phosphate group. Metal ions at both sites have been suggested to participate directly in the chemical step (Peracchi et al. 1997; Wang et al. 1999). By contrast, other biophysical techniques such as single molecule FRET and isothermal calorimetry support delocalized Mg\(^{2+}\) binding (Rueda et al. 2003; Mikulecky et al. 2004). By comparing the pH-rate profiles for Mg\(^{2+}\) and several divalent transition metal ions, we have proposed that divalent metal ions may support high rates of catalysis by perturbing the proton transfer character of guanosine nucleobases in the active site, perhaps through direct, inner-sphere coordination with the guanosines (Roychowdhury-Saha and Burke 2006).

Insight into the role(s) of divalent ions in ribozyme catalysis can be gained by studying catalysis in solutions with other cations used in place of the divalent metal ions. Modest catalytic activity has also been observed for minimal hammerhead ribozymes in monovalent ions, polyamines, and exchange-inert cations (Murray et al. 1998; Curtis and Bartel 2001; O’Rear et al. 2001). Activity in monovalent cations correlated in log-linear fashion with ionic radius and approached that observed in Mg\(^{2+}\), suggesting that hydrated divalent ions do not act directly as a general base in the reaction (Curtis and Bartel 2001). However, a comparison of activities among a series of ribozyme variants in 4 M Li\(^{+}\) and in 10 mM Mg\(^{2+}\) suggested that important interactions were absent in the monovalent reactions (O’Rear et al. 2001). Polyamines are abundant inside cells, where they influence gene expression, DNA condensation, and RNA folding (Koculi et al. 2004). In ribozyme catalysis, they are primarily used to augment self-cleavage or substrate cleavage in the presence of divalent cations (Dahm and Uhlenbeck 1991; Suh et al. 1993; Earnshaw and Gait 1998; Akashi et al. 2002). Very slow cleavage was reported for a minimal hammerhead ribozyme in 0.5 mM spermine (Dahm and Uhlenbeck 1991). Cobalt hexammine (CoHex, Co\([\text{NH}_3]_6^{3+}\)) is a structural analog of fully hexahydrated Mg\(^{2+}\) (Mg\([\text{H}_2\text{O}]_6^{2+}\)) and—to a lesser degree—of partially dehydrated Mg\(^{2+}\) (Mg\([\text{H}_2\text{O}]_3^{2+}\)); thus, insight into the role of Mg\(^{2+}\) can be obtained from an analysis of the effects of substituting CoHex in place of Mg\(^{2+}\). The interactions of CoHex with RNA have been well documented in pseudoknots (Gonzalez and Tinoco 1999; Nixon et al. 1999), G–U tandem repeats (Kief and Tinoco 1997), stem–loop structures (Rudisser and Tinoco 2000), and helices (Butcher et al. 2000). In these contexts, CoHex exhibits delocalized, outer-sphere contacts with RNA similar to those provided by Mg\([\text{H}_2\text{O}]_6^{2+}\). In contrast to water molecules bound to Mg\(^{2+}\), the NH\(_3\) ligands of CoHex exchange very slowly (\(k\text{exchange} \sim 10^{-10}\ \text{s}^{-1}\)) (Basolo and Pearson 1988), effectively preventing access to inner-sphere coordination. The consequence of cobalt hexammine binding on ribozyme activity has been investigated in several ribozymes. In the hairpin ribozyme (Hampel and Cowan 1997) and in an in vitro selected acyl-transferase ribozyme (Suga et al. 1998), CoHex promotes efficient cleavage in the absence of divalent metal ions. In the Neurospora crassa VS ribozyme, it enhances activity in the presence of Mg\(^{2+}\) but cannot support catalysis as the sole metal ion (Maguire and Collins 2001). In the HDV ribozyme (Nakano et al. 2000) and in a minimal hammerhead ribozyme (Horton and DeRose 2000), CoHex inhibits activity in the presence of other divalent ions, suggesting a competition for binding. Previous studies with minimal hammerhead ribozymes showed very slow cleavage activity (~0.0071 min\(^{-1}\)) at high (>100 mM) cobalt hexammine concentrations (Curtis and Bartel 2001).

Artificial tertiary stabilizing motifs that function in the context of trans-cleaving versions of two natural cis-cleaving molecules were recently identified using a selection-amplification approach (Saksmerprome et al. 2004). A hammerhead ribozyme, designated RzB (Fig. 1), carrying one of these artificial tertiary motifs exhibits exceptionally high rates in the range of 100–3000 min\(^{-1}\) in 1 mM divalent cations (Roychowdhury-Saha and Burke 2006). We observed an increase in thermal stability upon tertiary stabilization both for ribozyme RzB (Saksmerprome et al. 2004) and for the larger ribozyme RzB0 (Roychowdhury-Saha and Burke 2006) (see FIGURE 1). Secondary structure of RzB annealed to an 18-nt substrate (Saksmerprome et al. 2004; Roychowdhury-Saha and Burke 2006). Nucleotides of the catalytic core are shown in red, nucleotides of the substrate strand are shown in blue, and the nucleotides of the tertiary stabilization motifs in Loop II and Bulge 1 of the ribozyme strand are shown in purple. RzB0 sequence lacks the “UAA” loop.

FIGURE 1. Secondary structure of RzB annealed to an 18-nt substrate (Saksmerprome et al. 2004; Roychowdhury-Saha and Burke 2006). Nucleotides of the catalytic core are shown in red, nucleotides of the substrate strand are shown in blue, and the nucleotides of the tertiary stabilization motifs in Loop II and Bulge 1 of the ribozyme strand are shown in purple. RzB0 sequence lacks the “UAA” loop.
Hammerhead catalysis in non-divalent-metal cations

RESULTS

Spermidine and Na⁺ promote RzB catalysis in the absence of divalent cations

Single-turnover trans-cleavage of a 13-nt RNA substrate was monitored for hammerhead ribozyme RzB as a function of spermidine. The observed rate of cleavage increased with spermidine at low concentrations and saturated at ∼10 mM spermidine with k_{max} = 0.025 ± 0.001 min⁻¹ (Fig. 2A). Previous studies of catalysis by a minimal hammerhead ribozyme in polyamines (Dahm and Uhlenbeck 1991) reported a rate of 0.00024 min⁻¹ in 0.5 mM spermine (a tetra-amine) at 25°C, which is ∼100-fold slower than the rate we observe here in spermidine (a tri-amine). The tertiary stabilization afforded by the peripheral sequence elements likely accounts for these differences in polyamine utilization. This interpretation is supported by the absence of cleavage activity in the control ribozyme lacking tertiary stabilization (ribozyme RzB0), even at 50 mM spermidine after 2 h (data not shown). Our results support the existence of an alternate folding pathway or catalytic mechanism used by the ribozyme in the presence of nondivalent cations (especially CoHex) that is distinct from that promoted by divalent cations.
The maximal extent of cleavage by RzB at long incubation times is similar in 5–20 mM spermidine (70%), 1 M Na⁺ (80%), and 1 mM Mg²⁺ (>90%) (Fig. 2). Thus, the RzB ribozyme is capable of accessing a catalytically competent conformation in each of these cations. However, the maximal cleavage rate observed in spermidine or Na⁺ is 10,000-fold lower than the maximal rate of 357 ± 23 min⁻¹ in Mg²⁺ (10 mM) under otherwise identical conditions (Roychowdhury-Saha and Burke 2006). This dramatic difference in rate may potentially reflect a different mechanism or rate-limiting step. Multichannel reaction pathways involving divalent-dependent and divalent-independent cleavage mechanisms have been noted previously for the HDV ribozyme and for a minimal hammerhead ribozyme (Nakano et al. 2001, 2003; Zhou et al. 2002). Analogous multiple pathways for hammerhead ribozymes may reflect different local architectures of the active site—such as the presence or absence of specific coordination of a metal ion to the scissile phosphate or to a nucleobase—even if the global folded structures of the ribozyme–substrate complex are similar. Such scenarios could affect the energetics of the cleavage reaction and result in dissimilar rates, particularly if only a subset of the cations can play a role that extends significantly beyond gross structural stabilization (see Discussion).

**CoHex promotes RzB catalysis**

Both inner- and outer-sphere mechanisms have been proposed for metal-ion-mediated catalysis by hammerhead ribozymes. To discriminate between these two possibilities, substrate cleavage by RzB was monitored in the presence of CoHex. The kinetic profile has two distinct phases, with a rapid burst that is complete within the first 10 sec, followed by a slower phase (Fig. 3A,B). In previous work, we (Roychowdhury-Saha and Burke 2006) and others (Canny et al. 2004; Zamel et al. 2004) have used a quench-flow strategy to measure rapid ribozyme cleavage mediated by Mg²⁺ or transition metal ions. A critical aspect of these experiments is the ability to quench the reaction on millisecond time scales by sequestering the metal ions with EDTA. This approach is not available for assessing RzB cleavage in CoHex, as this exchange-inert metal ion complex cannot be sequestered by EDTA. An alternative quench strategy employing strong denaturing conditions was explored (3 volumes of 99% formamide); however, this strategy was ineffective, as CoHex-promoted cleavage was faster than denaturation of the RzB active structure. Burst phase rates therefore represent lower-limit estimates, and a Hill coefficient for CoHex was not explicitly determined.

Both the rate of cleavage during the slow phase (assumed to reflect rearrangement from an inert to an active conformation) and the magnitude of the burst fraction (cleavage occurring during the first 10 sec, assumed to reflect the fraction that is prefolded in a catalytically competent conformation) are stimulated by increasing concentrations of CoHex (Fig. 3A,B). In 20 mM CoHex, the burst phase rate (≥8.9 ± 0.9 min⁻¹) is higher than any previously reported for a hammerhead ribozyme in the presence of CoHex. Cleavage rates during the slow phase ranged from 0.012 ± 0.002 min⁻¹ (0.1 mM CoHex) to 0.149 ± 0.027 min⁻¹ (20 mM CoHex). The low end of this range is similar to values obtained previously (0.009 min⁻¹) for the extended *Schistosoma* hammerhead in 0.1 mM CoHex (Osborne et al. 2005). RzB0 does not show detectable cleavage after 2 h even at CoHex concentrations as high as 20 mM (data not shown), similar to the nonreactivity or poor reactivity observed previously for minimal hammerhead ribozymes (Curtis and Bartel 2001; Osborne et al. 2005).

**Competition between Mg²⁺ and CoHex**

To the extent that Co(NH₃)₆³⁺ acts as a true structural analog of Mg(H₂O)₆²⁺ (similar geometry and size), both ions together should have additive effects, whereas if they promote catalysis via different mechanistic pathways or...
promote alternative folds, they are expected to exhibit competitive behavior. Therefore, competition assays were performed to address the potential synergy or antagonism in the cleavage mechanisms stimulated by Mg$^{2+}$ versus CoHex (Fig. 4; Table 1). Cleavage kinetics in a constant background of 0.2 mM Mg$^{2+}$ were monitored in various concentrations of CoHex (Fig. 4A). The initial rate decreased as CoHex concentration was increased to \( \approx 1.0 \) mM (Fig. 4B), but at still higher concentrations, cleavage increased markedly and exhibited burst kinetics (Fig. 4A,C). Throughout this series of reactions, increases in the CoHex concentration yielded a concomitant decrease in the fraction cleaved at equilibrium (\( F_N \)) (Fig. 4A). Time courses at 0.3 and 0.5 mM constant Mg$^{2+}$ showed similar decreases in \( F_N \) with increasing CoHex (data not shown). In contrast, when no Mg$^{2+}$ was present, increasing the CoHex concentration increased both the burst phase population and the final equilibrium fraction, \( F_N \) (cf. Fig. 3). Both of these results suggest differences in the mechanistic details of catalysis by ribozyme RzB in the presence of CoHex versus in the presence of Mg$^{2+}$ (see Discussion).

### DISCUSSION

#### Multichannel catalysis by ribozyme RzB

The catalytic activities reported here for hammerhead ribozyme RzB in the presence of sodium, spermidine, and CoHex confirm and extend previous findings of divalent-ion-independent hammerhead ribozyme catalysis. For ribozyme RzB, the difference between reaction rates observed in divalent ions (Roychowdhury-Saha and Burke 2006) and those observed in Na$^+$ or spermidine (Fig. 2) is likely due to tertiary stabilization in RzB, which could then result in differing kinetic pathways. Previous modeling of minimal hammerhead ribozyme kinetics suggested a multichannel reaction mechanism with Mg$^{2+}$-dependent, Li$^+$-dependent, and cooperative pathways (Takagi et al. 2004). In those studies, the reaction in divalent ions was \( \approx 20 \)-fold faster than in monovalent ions (Curtis and Bartel 2001). Previous studies of the Mg$^{2+}$ dependence of substrate cleavage by RzB yielded a Hill coefficient (for catalysis) of 2.0 ± 0.3 (Roychowdhury-Saha and Burke 2006), suggesting a requirement for two metal ions to access the catalytic

<table>
<thead>
<tr>
<th>[CoHex], mM</th>
<th>No Mg$^{2+}$</th>
<th>0.2 mM Mg$^{2+}$</th>
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<tr>
<td>0</td>
<td>ND</td>
<td>57</td>
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<tr>
<td>0.1</td>
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<td>0.5</td>
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<td>20</td>
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(ND) Not determined.
The presence of Mg2+ alone (Roychowdhury-Saha and Burke 2005) RNA, Vol. 13, No. 6 distinctive metal ion dependencies (Alam et al. 2005). In addition, mutations near the hairpin catalytic core induced conformational mobility that may be ion dependent. In cross-linking is on a path to catalysis, these results suggest a mechanistic detail from the pathway promoted by Mg(H2O)n2+. First, the maximal rate attained by RzB in the presence of Mg2+ alone (Roychowdhury-Saha and Burke 2006) is as much as 40-fold higher than that attained in CoHex (357 ± 23 versus 8.9 ± 0.9 min⁻¹). A caveat here is that because the CoHex rate is a lower-limit value, the actual difference in initial rate could be significantly less. Second, the calculated values of Fₙ at 55 ± 5% for reactions carried out in the presence of CoHex (Fig. 3A) are significantly lower than those seen in Mg2+ or other divalent ions at >90% or in sodium or spermidine at 70%–80% (Fig. 2B; Roychowdhury-Saha and Burke 2006). Thus, the ribozyme is less able to access a catalytically competent conformation in CoHex even at long times. Third, CoHex and Mg2+ compete for binding the ribozyme. Cleavage rates in a constant Mg2+ background initially decrease with increasing CoHex concentrations as CoHex displaces Mg2+ and interferes with Mg2+-mediated catalysis. Still higher concentrations lead to increases in the initial rates and final cleavage extents and to burst kinetics, but the rates and yields do not return to those observed at elevated CoHex in the absence of Mg2+ (Table 1). Thus, CoHex interferes with Mg2+-mediated catalysis and Mg2+ interferes with CoHex-mediated catalysis.

We interpret these data to indicate that hammerhead ribozyme RzB can fold into a Mg2+-dependent or a Mg2+-independent active state in the presence of either metal ion alone, but that simultaneous binding by both metal ions creates an intermediate state with little or no catalytic activity (Fig. 5). Competition between CoHex and Mg2+ ions, with accompanying structural changes, has also been observed in a minimal hammerhead ribozyme (Horton and DeRose 2000).

Mechanistic model for reduced activity in the absence of divalent ions

The “alternate” reaction pathways promoted by Na+, spermidine, and CoHex could be either structural or mechanistic in nature. For example, photo-cross-linking studies (Lambert et al. 2006a,b) suggest a different active site geometry in the hairpin ribozyme in CoHex than in the structures proposed from crystallographic studies in Ca2+ (Rupert and Ferre-D’Amare 2001) or in CoHex (Alam et al. 2005). To the extent that the conformation captured by cross-linking is on a path to catalysis, these results suggest a conformational mobility that may be ion dependent. In addition, mutations near the hairpin catalytic core induced distinctive metal ion dependencies (Alam et al. 2005). In contrast, Neurospora VS ribozyme shows similar conformational changes in both Mg2+ and CoHex during chemical modification structure probing assays, although Mg2+ is more effective than CoHex in inducing important structural changes (Maguire and Collins 2001).

For the present work an attractive alternative is that the “alternate pathways” involve removal of a specific catalytic mechanism that is available to Mg2+ and to transition metal ions but not to monovalent ions, polyamines, or exchange-inert cations. There is significant debate regarding the functional and structural roles of Mg2+ in hammerhead ribozyme catalysis. Magnesium can act as a Lewis acid via inner-sphere contacts or by any of several outer-sphere interactions, such as hydrogen bonding, diffuse Coulombic interactions, or proton transfer. Its potential role in proton transfer has been called into question by recent studies in which simultaneous substitution of G8 and G12 with 2,6-diaminopurine (diAP; pKa 5.1 versus pKa 9.6 for G) in a minimal hammerhead ribozyme yielded a bell-shaped pH-rate profile, implicating these two nucleobases in both proton transfer steps (Han and Burke 2005). Furthermore, our recent analysis of the pH dependence of substrate cleavage by RzB in various metal ions suggests that the metal ion hydrate is not acting as the general base for the reaction (Roychowdhury-Saha and Burke 2006). However, the pKa of the general acid correlates with the pKa of the metal ion hydrate, displaced to lower values by 1.0–1.7 units (Roychowdhury-Saha and Burke 2006), suggesting that the metal ion may perturb one of the guanosine’s pKa by a direct (inner-sphere) coordination of the metal ion to the nucleobase or by proximity of the metal ion’s positive charge. The slower catalysis observed in Na+, spermidine, and cobalt hexammine may therefore stem from the
decreased ability for general acid protonation of the 5′ oxyanion leaving group due to the inability of these cations to form strong inner-sphere coordination with an active site guanosine.

MATERIALS AND METHODS

Quantification of the ribozyme reaction

Kinetic measurements were performed under ribozyme-saturated, single-turnover conditions with two RNA constructs (RzB and RzB0), each of which was assembled from separate ribozyme and substrate strands. This design ensures that the conversion of the ribozyme–substrate complex to the ribozyme–product complex could be monitored without the complications arising from kinetic processes associated with annealing of the complex. Ribozyme strands were synthesized by in vitro transcription using T7 RNA polymerase from the Epicentre Ampliscribe kit following the supplier’s protocol. A chemically synthesized substrate strand (Integrated DNA Technologies) was 5′ radiolabeled using [γ-32P]ATP and polynucleotide kinase. All cleavage reactions were performed at 37°C, pH 7.4. Reactions were carried out with a 2 μM ribozyme strand and a 50 nM substrate strand in 50 mM Tris-HCl and 10 μM EDTA, as described previously (Saksmerprome et al. 2004). Briefly, each reaction was initiated by adding 5 μL of a cation solution to 70 μL of preannealed ribozyme–substrate solution at 37°C, pH 7.4. Aliquots (5 μL) were withdrawn at each time point and quenched with 1–2 volumes of 90% formamide, 50 mM EDTA, and 0.005% each of xylene cyanol and bromophenol blue. Quenched samples were stored at −80°C prior to denaturing (8 M urea) polyacrylamide gel electrophoresis. Metal ion competition experiments were initiated by addition of 5 μL of a mixture of MgCl2 and [Co(NH3)6]Cl2 to 70 μL of preannealed ribozyme–substrate solution at 37°C, pH 7.4. The extent of cleavage in each reaction was quantified using ImageQuant software (Molecular Dynamics). The fraction cleaved (Ft) at time t was fit using KaleidaGraph 3.5 either to a single-exponential equation:

\[ F_t = F_0 + (F_{\infty} - F_0) \left(1 - \exp \left(-\frac{t}{k_{obs,1}}\right)\right) \]  

(1)

or to a biexponential equation:

\[ F_t = F_0 + (F_{\infty} - F_0) \left\{ \left(1 - \alpha \right) \exp \left(-\frac{t}{k_{obs,1} \cdot \alpha}\right) - (1 - \alpha) \exp \left(-\frac{t}{k_{obs,2} \cdot (1-\alpha)}\right) \right\} \]  

(2)

where \( F_0 \) is the zero point correction, \( F_{\infty} \) is the estimated plateau value at infinite time, and \( \alpha \) and \( (1 - \alpha) \) are the fractions of the cleaved population that have rate constants of \( k_{obs,1} \) and \( k_{obs,2} \), respectively. Reported errors reflect uncertainties of the KaleidaGraph fit of the data to these equations.

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REFERENCES


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