

Structure of a Plant Cell Wall Fragment Complexed to Pectate Lyase C

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The three-dimensional structure of a complex between the pectate lyase C (PelC) R218K mutant and a plant cell wall fragment has been determined by x-ray diffraction techniques to a resolution of 2.2 Å and refined to a crystallographic *R* factor of 18.6%. The oligosaccharide substrate, α -D-GalpA-([1→4]- α -D-GalpA)₃-(1→4)-D-GalpA, is composed of five galacturonopyranose units (D-GalpA) linked by α -(1→4) glycosidic bonds. PelC is secreted by the plant pathogen *Erwinia chrysanthemi* and degrades the pectate component of plant cell walls in soft rot diseases. The substrate has been trapped in crystals by using the inactive R218K mutant. Four of the five saccharide units of the substrate are well ordered and represent an atomic view of the pectate component in plant cell walls. The conformation of the pectate fragment is a mix of 2₁ and 3₁ right-handed helices. The substrate binds in a cleft, interacting primarily with positively charged groups: either lysine or arginine amino acids on PelC or the four Ca²⁺ ions found in the complex. The observed protein–oligosaccharide interactions provide a functional explanation for many of the invariant and conserved amino acids in the pectate lyase family of proteins. Because the R218K PelC–galacturonopentaose complex represents an intermediate in the reaction pathway, the structure also reveals important details regarding the enzymatic mechanism. Notably, the results suggest that an arginine, which is invariant in the pectate lyase superfamily, is the amino acid that initiates proton abstraction during the β elimination cleavage of polygalacturonic acid.

INTRODUCTION

Pectate lyases are depolymerizing enzymes that degrade plant cell walls, causing tissue maceration and death. The enzymes normally are secreted by phytopathogenic organisms and are known to be the primary virulence agents in soft rot diseases caused by *Erwinia* spp (Collmer and Keen, 1986; Kotoujansky, 1987; Barras et al., 1994). In the latter organisms, the enzymes exist as multiple, independently regulated isozymes that share amino acid sequence identity ranging from 27 to 80%.

Pectate lyases share sequence similarities with fungal pectin lyases, plant pollen proteins, and plant style proteins (Henrissat et al., 1996). The three-dimensional structures of

five members of the superfamily have been determined and include *Erwinia chrysanthemi* pectate lyase C (PelC) (Yoder et al., 1993; Yoder and Journak, 1995), *E. chrysanthemi* pectate lyase E (PelE) (Lietzke et al., 1994), *Bacillus subtilis* pectate lyase (B. *subtilis* Pel) (Pickersgill et al., 1994), *Aspergillus niger* pectin lyase A (PLA) (Mayans et al., 1997), and *A. niger* pectin lyase B (PLB) (Vitali et al., 1998). All share a similar but an unusual structural motif, termed the parallel β helix, in which the β strands are folded into a large, right-handed coil. The enzyme structures differ in the size and conformation of the loops that protrude from the parallel β helix core. As deduced from sequence similarity and site-directed mutagenesis studies, the protruding loops on one side of the parallel β helix form the pectolytic active site (Kita et al., 1996). The structural differences of the loops are believed to be related to subtle differences in the enzymatic and maceration properties of the proteins.

Pectate lyases catalyze the cleavage of pectate, the de-esterified product of pectin, which is the major component that maintains the structural integrity of cell walls in higher plants

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(Carpita and Gibeaut, 1993). The pectate backbone is composed of blocks of polygalacturonic acid (PGA), which is a helical homopolymer of D-galacturonic acid (GalpA) units linked by α -(1 \rightarrow 4) glycosidic bonds. The blocks of PGA are separated by stretches in which (1 \rightarrow 2)- α -L-rhamnose residues alternate with GalpA (McNaught, 1997). Blocks of PGA may contain as many as 200 GalpA units and span 100 nm (Thibault et al., 1993). Cations are necessary to neutralize PGA in solution and, as a consequence, influence its structure.

In the presence of Ca^{2+} , PGA assumes a 2_1 helical conformation in dilute polymer concentrations (Morris et al., 1982; Powell et al., 1982) and a 3_1 helix at high concentrations in either a gel or solid form (Walkinshaw and Arnott, 1981a, 1981b). Because the PGA concentration in the plant cell wall upon demethylation of pectin lies near the critical conformational transition point, considerable speculation exists as to the in situ structure of PGA. A popular view is the "eggbox model" in which Ca^{2+} ions cross-link the uronic acid moieties of neighboring antiparallel chains of PGA together. Although the eggbox model generally is depicted with PGA in a right-handed 2_1 helical conformation, the original literature suggests that cross-linking between Ca^{2+} and PGA in a right- or left-handed 3_1 helical conformation is feasible as well (Grant et al., 1973; Kohn, 1975).

The results of a recent nuclear magnetic resonance study suggest that the Ca^{2+} -PGA complex in the plant cell wall is much more complex than the simple eggbox model. This complex contains both 2_1 and 3_1 helices of PGA as well as intermediate conformational states (Jarvis and Apperley, 1995). Nuclear magnetic resonance, molecular modeling, and molecular dynamic analyses of pectic disaccharides and trisaccharides also have reported that PGA has both 3_1 and 2_1 helical conformations (Hricovini et al., 1991; DiNola et al., 1994; Gouvion et al., 1994). Disaccharide hydration and sodium salt formation may shift the predicted PGA helical conformation from 3_1 to 2_1 (DiNola et al., 1994; Gouvion et al., 1994; Catoire et al., 1997).

All proteins in the pectate lyase superfamily are believed to share a similar enzymatic mechanism, but the catalytic roles of the amino acids in the active site region have not been identified. For reasons of technical convenience, recent studies have focused on PelC. The enzyme randomly cleaves PGA by a β elimination mechanism, generating primarily a trimer end product with a 4,5-unsaturated bond in the galacturonosyl residue (α -L-4-eno-threohexosylpyranosyluronic acid [α -L-4-en-thrHexpA]) at the nonreducing end (Preston et al., 1992). PelC has an in vitro pH optimum of 9.5 and requires Ca^{2+} for pectolytic activity. Structural studies

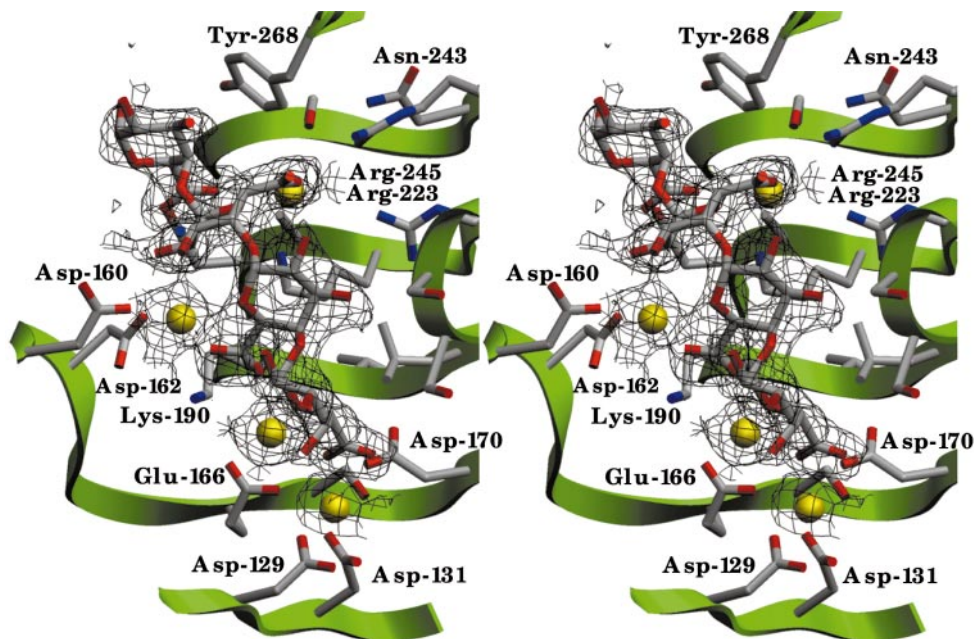


Figure 1. Stereoview of the Ca^{2+} ions and TetraGalpA Superimposed upon the Simulated Annealed OMIT Electron Density Map of the PelC R218K-Substrate Complex Contoured at 1.0σ .

The Ca^{2+} ions are represented by yellow spheres. TetraGalpA as well as the interacting amino acids are represented by rods by using the International Union of Pure and Applied Chemistry coloring code: carbon atoms are gray; oxygen atoms, red; and nitrogen atoms, blue. The R218K backbone is represented by green ribbons. Individual amino acids that are shown are labeled at the α -carbon.

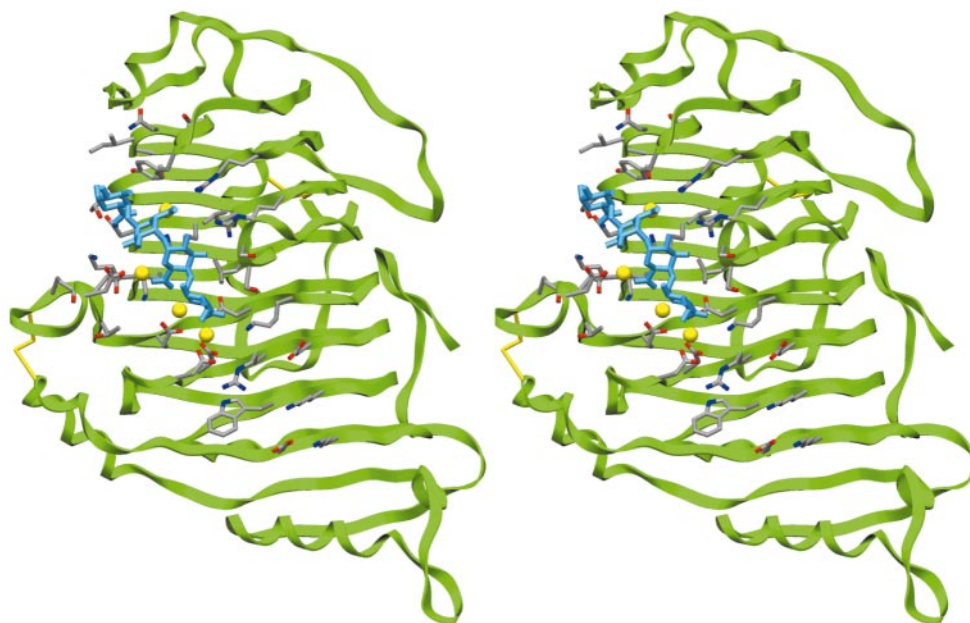


Figure 2. Stereoview of the PelC R218K-(Ca²⁺)₃₋₄-PentaGalpA Complex.

The view and color scheme are the same as given in Figure 1, except that the tetraGalpA substrate is illustrated in cyan, the entire protein backbone is shown, and the individual amino acids are not labeled. The disulfide bonds are illustrated as yellow rods.

have shown that Ca²⁺ is bound to the enzyme at a location that was first suggested in a PelC-Lu³⁺ complex (Yoder et al., 1993) and later confirmed by structural studies of a *B. subtilis* Pel-Ca²⁺ complex (Pickersgill et al., 1994). The role of Ca²⁺ has not been established. In the β elimination reaction, the reaction is initiated by proton abstraction from C-5 of the galacturonosyl residue on the reducing side of the glycosidic scissile bond. The group or groups that initiate proton abstraction and transfer the proton to the glycosidic oxygen have not been identified. Potential candidates include two invariant amino acids in the superfamily, Asp-131 and Arg-218 in PelC nomenclature, as well as four amino acids, Glu-166, Asp-170, Lys-190, and Arg-223, which are invariant within the pectate lyase subfamily (Henrissat et al., 1996). Site-specific mutations at the latter PelC positions abolish pectolytic as well as maceration activity (Kita et al., 1996). Notably, the pectolytic region is devoid of conserved histidine, serine, or tyrosine residues, which frequently are implicated in β elimination enzymatic mechanisms with a lower pH optimum. In this study, we have taken advantage of the impaired catalytic properties of one PelC mutant, R218K, to form a stable substrate-enzyme complex that could be studied by x-ray diffraction techniques. The results provide an atomic view of a pectate fragment, α -D-GalpA-([1 \rightarrow 4]- α -D-GalpA)₃-(1 \rightarrow 4)-D-GalpA (pentaGalpA), and the identification of the key amino acids involved in oligosaccharide

binding. In addition, the results provide tentative identification of the amino acid that initiates proton abstraction.

RESULTS

Conformation of the PentaGalpA Substrate

Four of the five GalpA units of the substrate used in the crystal diffraction experiment are visible as strong, well-ordered electron density in difference Fourier maps, as shown in Figure 1. As shown in Figure 2, the well-ordered GalpA units interact with PelC in a groove encompassing the previously identified Ca²⁺ binding site on the protein, now termed the ⁴Ca²⁺ site. The orientation of the tetraGalpA fragment is unambiguous. The reducing end, GalpA¹, is located at the protein-solvent border, and the nonreducing end, GalpA⁴, lies near ⁴Ca²⁺. Additional electron density, corresponding to a fifth GalpA unit, is found at the nonreducing end of the tetraGalpA fragment but is disordered and cannot be modeled.

Each of the well-ordered GalpA rings refines to the conventional chair conformation, with all bond distances and angles consistent with single bonds. The pectate fragment

Table 1. Bond Angles (τ) and Torsional Rotations (ϕ and ψ) about Glycosidic Bonds in the Refined Structures of the PelC-(Ca²⁺)₃₋₄-PentaGalpA Complexes^a

Glycosidic Bond	τ^b	ϕ^c	ψ^d
3 ₁ Helix ^e	117.0°	80.0°	89.0°
2 ₁ Helix ^f	117.0°	80.0°	161.0°
GalpA ¹ -GalpA ²	117.0°	54.0°	90.0°
GalpA ² -GalpA ³	116.6°	117.0°	157.0°
GalpA ³ -GalpA ⁴	120.4°	73.0°	51.0°

^aTorsional angles were determined by looking from the nonreducing end side (with prime) down the bond of interest to the reducing end side and determining the angle of rotation created from the planes of O-5'-C-1'-O-4 and C-1'-O-4-C-4 for ϕ and C-1'-O-4-C-4 and O-4-C-4-C-5 for ψ . A *cis* configuration is taken to be 0°, and a *trans* configuration is taken to be 180°. A negative sign is a rotation from *cis* in a counterclockwise direction, and a positive sign is a rotation from *cis* in a clockwise direction.

^b τ , the C₁-O-C₄' bond angle.

^c ϕ , the torsional rotation about the C₁-O bond.

^d ψ , the torsional rotation about the O-C₄' bond.

^eThe values for the 3₁ helix are those determined by the program O (Jones et al., 1991) from the model published by Walkinshaw and Arnott (1981b).

^fThe values for the 2₁ helix are those determined by the program O for a galacturonic acid model constructed using the parameters for alginic acid by Atkins et al. (1973).

folds into an unbent right-handed helical conformation, with the observed helical angles compared with the idealized 2₁ and 3₁ helices in Table 1. Two of the three glycosidic bonds have ψ torsional angles that approximate the 3₁ helix observed in fiber diffraction studies of PGA-Ca²⁺ gels (Walkinshaw and Arnott, 1981a, 1981b). One of the glycosidic bonds, between GalpA² and GalpA³, has a ψ torsional angle similar to a 2₁ helix. Consequently, the overall appearance of the pectate fragment conformation is of a 3₁ helix, but with the middle segment distorted into 2₁ helix, as illustrated in Figures 3A and 3B. Given the greater number of GalpA³ contacts, as listed in Table 2, the deviation from the 3₁ helical conformation is probably a result of specific interactions with the enzyme. The deviation is not likely due to other effects, such as pectate concentration, hydration, or cation type, which are postulated to cause the transition between 2₁ and 3₁ helical conformations of pectate. If the R218K-(Ca²⁺)₃₋₄-GalpA₅ structures are representative of interactions that occur within the plant cell wall, then endogenous proteins also are likely to distort the PGA conformation from any helical states observed under in vitro conditions.

Coordination of Ca²⁺ Ions

In wild-type PelC, a Ca²⁺ ion coordinates to seven ligands, including two water molecules, both carboxyl oxygens of Asp-

131, and one carboxyl oxygen from each of Asp-129, Glu-166, and Asp-170. In the R218K complex with pentaGalpA, the equivalent ⁴Ca²⁺ coordinates to the same groups, with a single exception—a carboxyl oxygen from GalpA⁴ replaces one of the water molecules. In addition to ⁴Ca²⁺, three additional Ca²⁺ ions have been identified. Two of the additional Ca²⁺ ions, ²Ca²⁺ and ³Ca²⁺, are fully occupied, and the third, ¹Ca²⁺, has a partial occupancy of ~50%. Each Ca²⁺ ion bridges the carboxyl group of each GalpA unit to the protein. In addition, ²Ca²⁺ and ³Ca²⁺ link the uronic acid moieties of GalpA², GalpA³, and GalpA⁴. The coordination around each Ca²⁺ ion is listed in Table 3. The observed Ca²⁺ positions are very different from the interstrand Ca²⁺ ions

Table 2. Atomic Distances of 3.0 Å or Less between the Oxygen Atoms of TetraGalpA and Amino Acids, Ca²⁺ Ions, or Water Molecules

GalpA Atoms ^a	Interacting Atoms ^b	Distance (Å)
GalpA ¹		
Ring interaction	Tyr-268	
O-6A	GA ¹ Wat ¹	2.9
O-6B	¹ Ca ²⁺	2.9
GalpA ²		
O-2	Asp-162 O- δ 2	2.7
O-2	GA ² Wat ¹	2.7
O-3	² Ca ²⁺	2.6
O-5	¹ Ca ²⁺	2.9
O-6A	¹ Ca ²⁺	2.6
O-6A	Arg-245 NH-1	2.8
O-6B	Arg-245 NH-2	3.0
O-6B	GA ² Wat ²	2.8
GalpA ³		
O-2	Arg-223 NH-2	2.9
O-3	Ser-196 O	2.8
O-3	Arg-223 NH-1	2.9
O-5	² Ca ²⁺	2.5
O-5	² CaWat ²	3.0
O-6A	Lys-190 N- ζ	2.8
O-6A	² Ca ²⁺	2.3
O-6A	² CaWat ²	2.9
O-6B	³ Ca ²⁺	2.4
O-6B	³ CaWat ³	2.8
GalpA ⁴		
O-2	² CaWat ²	2.9
O-3	GA ⁴ Wat ¹	2.8
O-4	GA ⁴ Wat ²	2.7
O-4	Ser-308 O	2.8
O-5	³ Ca ²⁺	2.4
O-6A	⁴ Ca ²⁺	2.5
O-6A	³ Ca ²⁺	2.4
O-6A	GA ⁴ Wat ²	2.9
O-6B	⁴ CaWat ¹	2.6

^aThe positions of the atoms are indicated in Figure 4.

^bXWat^Y refers to the Y water molecule associated with the X GalpA unit. ^ZCa²⁺ refers to the Z position of the Ca²⁺ ion as defined in Figure 4.

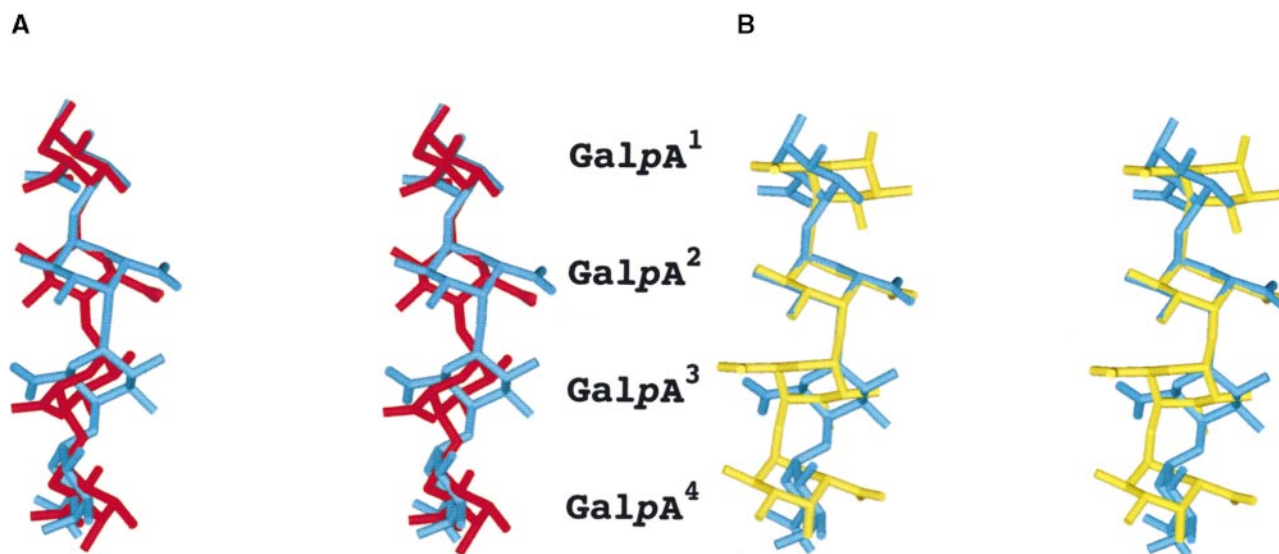


Figure 3. Stereoview of the TetraGalpA Structure Superimposed upon Modeled Right-Hand 2_1 and 3_1 OligoGalpA Helices.

The tetraGalpA structure determined at 2.2 Å is shown in both (A) and (B) as cyan rods.

(A) The modeled 3_1 oligoGalpA helix (Walkinshaw and Arnott, 1981a) is shown in red.

(B) The modeled 2_1 oligoGalpA helix (Atkins et al., 1973) is shown in yellow.

postulated to link PGA helices together (Walkinshaw and Arnott, 1981a, 1981b; Morris et al., 1982; Powell et al., 1982). In the present structure, the Ca^{2+} ions link not only the oligosaccharide to the protein but also adjacent uronic acid moieties within a single pectate strand.

Protein- Ca^{2+} -TetraGalpA Interactions

The protein- Ca^{2+} -tetraGalpA interactions are represented in Figure 4, and all relevant interatomic distances are summarized in Table 2. Electrostatic interactions dominate, with the negatively charged uronic acid moieties primarily interacting with positively charged groups: either lysine or arginine on PelC or the four Ca^{2+} ions found in the complex. The carboxyl oxygens of GalpA² and GalpA³ interact strongly with Arg-245 and Lys-190, respectively, whereas a carboxyl oxygen of GalpA⁴, at a distance of 3.2 Å from Lys-172, forms a weaker interaction. Lys-172 is highly conserved, and Lys-190 is invariant in the pectate lyases, but neither amino acid is found among the pectin lyases that bind a neutral methylated form of pectate. Arg-245 is conserved only among PelC subfamily members but not in the PelE subfamily whose members rapidly cleave the substrate to an unsaturated dimer. Several additional interactions between tetraGalpA and the protein were observed, but notably, the most specific ones involve GalpA³. Arg-223, another invariant amino acid in the pectate lyase subfamily, forms hydro-

gen bonds with the C-2 and C-3 hydroxyl groups of GalpA³, the orientation of which partially defines the galactose epimer. The C-3 hydroxyl group also forms a hydrogen bond with a nonconserved Ser-196. In GalpA², the C-2 hydroxyl interacts with Asp-162, and in GalpA¹, the ring forms a stacking interaction with Tyr-268. Both amino acids are conserved but only in the PelC subfamily. In addition to interactions with the protein and Ca^{2+} ions, the tetraGalpA segment is highly solvated, forming many hydrogen bonds with water molecules that increase in frequency from GalpA¹ to GalpA⁴. Collectively, the observed protein-tetraGalpA interactions provide a functional role for all invariant and conserved amino acids in the pectolytic region of the pectate lyases, except one, Arg-218.

Position of Scissile Bond

Crystals of wild-type PelC, which are isomorphous with R218K crystals, cleave pentaGalpA when diffused into crystals. Because the R218K mutant is catalytically inactive and a saturated tetraGalpA has been observed, the R218K- $(\text{Ca}^{2+})_{3-4}$ -GalpA₅ complexes represent a Michaelis complex in the reaction pathway. Can the scissile glycosidic bond be identified with certainty? PelC and subfamily members have been reported to cleave a pectate substrate, yielding a trimer as the primary unsaturated end product (68 to 72%; Preston et al., 1992). In the crystal structure, an unsaturated trimeric end product would result only if the scissile bond

Table 3. Ca²⁺-Coordinating Ligands^a in the PelC R218K-(Ca²⁺)₃₋₄-PentaGalpA Complexes

Ca ²⁺	Ligand	Distance (Å)
¹ Ca ²⁺ ^b	Lys-218 N-ζ	2.6
	GalpA ¹ O-6B	2.9
	GalpA ² O-5	2.9
	GalpA ² O-6A	2.6
² Ca ²⁺	¹ CaWat ¹	2.8
	Asp-160 O-δ2	2.3
	Asp-162 O-δ2	2.4
	GalpA ² O-3	2.6
	GalpA ³ O-5	2.5
	GalpA ³ O-6A	2.2
	² CaWat ¹	2.6
³ Ca ²⁺	² CaWat ²	2.2
	Glu-166 O-δ1	2.5
	Glu-166 O-δ2	2.5
	GalpA ³ O-6B	2.4
	GalpA ⁴ O-5	2.4
	GalpA ⁴ O-6A	2.4
	³ CaWat ¹	2.5
	³ CaWat ²	2.4
⁴ Ca ²⁺	³ CaWat ³	2.4
	Asp-129 O-δ1	2.3
	Asp-131 O-δ1	2.5
	Asp-131 O-δ2	2.4
	Glu-166 O-δ1	2.4
	Asp-170 O-δ2	2.4
	GalpA ⁴ O-6A	2.5
⁴ CaWat ¹	2.2	

^a^zCa²⁺ refers to the Z position of the Ca²⁺ ion as defined in Figure 4. ^xWat^y refers to the Y water molecule associated with either the X GalpA unit or the X Ca²⁺ ion. The labels for the oxygen atoms are defined in Figure 4.

^bThe observed electron density is best suited to a Ca²⁺ with a 50% occupancy rather than to a water molecule. The ¹Ca²⁺ ion is in the same location, relative to the uronic acid of GalpA¹, as are the other Ca²⁺ ions that coordinate GalpA units. However, unfavorable contact with the well-ordered and fully occupied lysine, Lys-218, was observed. It is not possible to determine whether ¹Ca²⁺ interacts with an unprotonated lysine at pH 9.5 in the crystals in 50% of the molecules.

occurred between GalpA³ and GalpA⁴. Moreover, only interactions with GalpA³ and GalpA⁴ involve highly conserved and invariant amino acids within the pectate lyase family. GalpA³ forms the most protein interactions, which appear to cause the greatest distortion from the 3₁ helical conformation of the tetraGalpA. In contrast, there are fewer interactions with GalpA¹ and GalpA², and all involve amino acids that are conserved only within the PelC subfamily.

To confirm the position of the scissile glycosidic bond, we investigated the enzymatic cleavage patterns of oligogalacturonates with different degrees of polymerization under optimized assay conditions for PelC. The composition of both the saturated and unsaturated end products was analyzed,

and the results, in Table 4, demonstrate that a pentaGalpA substrate has two observed modes of binding on PelC. The primary mode yields, as products, an unsaturated trimer (4-en-thrHexpA-[GalpA]₂) and a saturated dimer at a frequency of 71%. A secondary binding mode occurs at a 29% frequency, yielding an unsaturated dimer (4-en-thrHexpA-GalpA) and a saturated trimer. When reduced pentaGalpA, containing a galactonic acid at the reducing end, is used as the substrate, the cleavage pattern produces a reduced, unsaturated tetramer ([4-en-thrHexpA-GalpA]₂-L-Gal-onic, where L-Gal-onic refers to L-galactonic acid) and a saturated monomer at a 68% frequency. In addition, the cleavage pattern produces a reduced, unsaturated trimer (4-en-thrHexpA-GalpA-L-Gal-onic) and a saturated dimer at a 32% frequency. The new pattern is indicative of a shift toward the nonreducing end in the position of the scissile bond, because the galactonic acid unit now lies outside the enzyme in the primary binding mode. Because the galactonic acid unit is open rather than in a ring structure, the reduced saccharide cannot participate in the same interactions and occupy the GalpA¹ site on the protein-substrate complex. The only bond position that is consistent with the observed primary-mode cleavage patterns for the reduced and unreduced pentaGalpA substrate is that between GalpA³ and GalpA⁴ in the crystals of the protein-substrate complex.

DISCUSSION

The β elimination reaction in pectolytic cleavage is believed to involve three processes: neutralization of the carboxyl group adjacent to the scissile glycosidic bond, abstraction of the C-5 proton, and transfer of the proton to the glycosidic oxygen. In the R218K-(Ca²⁺)₃₋₄-pentaGalpA structures, the carboxyl group of GalpA³ is neutralized by interactions with ³Ca²⁺ and ²Ca²⁺ as well as by Lys-190, an invariant amino acid in the pectate lyase subfamily. Lys-190 also may serve an additional role, which is to partially protonate the carboxylic acid group, stabilizing an enolic intermediate as postulated and defined by Gerlt and colleagues (Gerlt et al., 1991; Gerlt and Gassman, 1992, 1993). Either or both effects serve to decrease the pK_a (the negative log of the dissociation constant) of the α proton at C-5, making it more susceptible to an attack by a base.

It is more difficult to definitively identify the group(s) responsible for proton abstraction and transfer. In our structure, there are no amino acids, water molecules, or Ca²⁺ ions within 3 to 4 Å of any C-5 atom or glycosidic oxygen for any GalpA unit. If the wild-type PelC structure is superimposed upon the R218K-substrate structure, as shown in Figure 5, there are minimal changes in the conformation of any side chain. However, one guanidinium nitrogen of the wild-type amino acid Arg-218 is positioned within 2.6 Å of C-5 of GalpA³, and the other nitrogen, at a distance of 2.7 Å, interacts with an oxygen of the carboxyl group. The latter in-

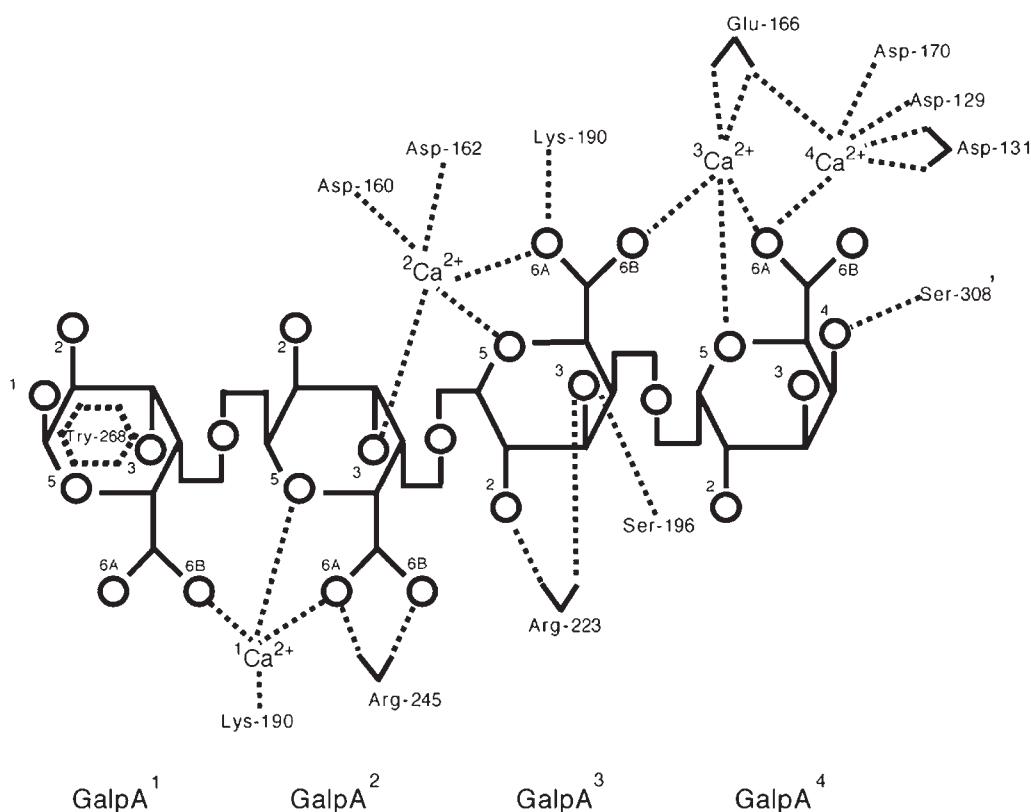


Figure 4. Schematic Representation of R218K and Ca^{2+} Ion Interactions with TetraGalpA at a Distance of $\leq 3.0 \text{ \AA}$.

GalpA¹ is the reducing saccharide, and GalpA⁴ is the nonreducing terminus. The interactions are designated with dotted lines, and the distances are given in Table 4. Oxygen atoms are represented by circles, with the corresponding number, and the carbon atoms are assumed at the intersection of bonds designated in boldface lines. Water molecules, which interact with tetraGalpA, are not shown but are listed in Table 4.

teraction is likely to be responsible for the lowered pK_a calculated for Arg-218. By using the MEAD program (Bashford and Gerwert, 1992), the calculated pK_a values for all PeIC arginine groups, except for Arg-218, fell within the range of 12.0 to 12.5. In contrast, the calculated pK_a value for Arg-218 is 9.5, approximately the same as the pH optimum of the reaction. It is highly unusual for an arginine to act as a general base during catalysis. However, as the H285R mutant of the acyl-acyl carrier protein thioesterase illustrates (Yuan et al., 1995), it is not impossible. The site-specific mutation of the catalytic histidine to an arginine shifts the enzymatic pH optimum from 8.5 to 12. In PeIC, the orientation of Arg-218 suggests a catalytic role, which is consistent with other known data, including the high pH optima for all pectate lyase-catalyzed reactions in vitro, the catalytic impairment of the R218K mutation, and the invariance of a comparable arginine in the pectate lyase superfamily.

In the structures presented in this study, no alternative atoms are close enough to the glycosidic oxygen between GalpA³ and GalpA⁴ to serve as a proton donor. When a partially flattened GalpA³ ring, expected during a β elimination

reaction, is modeled, a water molecule lies within 3 to 4 \AA of the glycosidic oxygen. The same water molecule, designated as $^3\text{CaWat}^2$, coordinates strongly to $^3\text{Ca}^{2+}$ and possibly is activated by the Ca^{2+} ion. Additional experiments are underway to test the novel enzymatic mechanism implied by the structural results.

In summary, the structures of the R218K- $(\text{Ca}^{2+})_{3-4}$ -penta-GalpA complexes provide an atomic view of a pectate component of the plant cell wall, revealing a right-handed, mixed 2_1 and 3_1 helical conformation for the observed tetraGalpA fragment and unanticipated Ca^{2+} positions. The complex represents a Michaelis complex in the reaction pathway, and the details have led to a possible catalytic mechanism, involving a novel role for an arginine as a C-5 proton abstractor. Moreover, the structure provides a functional explanation for all of the invariant and conserved residues in the pectolytic active site region of the pectate lyases. Unfortunately, the structure does not provide an explanation for another set of invariant residues, the vWIDH amino acid sequence, located in a second putative active site (Henrissat et al., 1996), but one that is too small to accommodate long

Table 4. End Product Analyses of PelC Cleavage of Oligogalacturonates

Oligomer Substrate ^a	Products ^b	Frequency (%)	Rate ($\mu\text{kat}/\text{mg}$)
(GalpA) ₃	GalpA + 4-en-thrHexpA-GalpA	100	0.17
(GalpA) ₄	GalpA + 4-en-thrHexpA-(GalpA) ₂	25	3 to 4
	(GalpA) ₂ + 4-en-thrHexpA-GalpA	75	
(GalpA) ₅	(GalpA) ₂ + 4-en-thrHexpA-(GalpA) ₂	71	10.1
	(GalpA) ₃ + 4-en-thrHexpA-GalpA	29	
(GalpA) ₆	(GalpA) ₂ + 4-en-thrHexpA-(GalpA) ₃	9	16.1
	(GalpA) ₃ + 4-en-thrHexpA-(GalpA) ₂	53	
	(GalpA) ₄ + 4-en-thrHexpA-GalpA	38	
(GalpA) ₇	(GalpA) ₂ + 4-en-thrHexpA-(GalpA) ₄	4	22.7
	(GalpA) ₃ + 4-en-thrHexpA-(GalpA) ₃	9	
	(GalpA) ₄ + 4-en-thrHexpA-(GalpA) ₂	55	
	(GalpA) ₅ + 4-en-thrHexpA-GalpA	32	
L-Gal-onic-(GalpA) ₂	No cleavage	0	
L-Gal-onic-(GalpA) ₃	GalpA + 4-en-thrHexpA-GalpA-L-Gal-onic	100	0.1
L-Gal-onic-(GalpA) ₄	(GalpA) ₂ + 4-en-thrHexpA-GalpA-L-Gal-onic	32	1.2
	GalpA + 4-en-thrHexpA-(GalpA) ₂ -L-Gal-onic	68	
L-Gal-onic-(GalpA) ₅	(GalpA) ₄ + 4-en-thrHexpA-L-Gal-onic	2	4.0
	(GalpA) ₃ + 4-en-thrHexpA-GalpA-L-Gal-onic	16	
	(GalpA) ₂ + 4-en-thrHexpA-(GalpA) ₂ -L-Gal-onic	82	
L-Gal-onic-(GalpA) ₆	(GalpA) ₅ + 4-en-thrHexpA-L-Gal-onic	8	4.0
	(GalpA) ₄ + 4-en-thrHexpA-GalpA-L-Gal-onic	21	
	(GalpA) ₃ + 4-en-thrHexpA-(GalpA) ₂ -L-Gal-onic	66	
	(GalpA) ₂ + 4-en-thrHexpA-(GalpA) ₃ -L-Gal-onic	5	

^a L-Gal-onic refers to L-galactonic acid or the reduced form of galacturonic acid.

^b 4-En-thrHexpA refers to α -L-eno-threohexosylpyranosyluronic acid or the 4,5-unsaturated form of galacturonic acid.

oligosaccharides. Despite the solvent accessibility in the crystals, no GalpA units are found near the vWIDH region, eliminating the possibility that the invariant amino acids are involved in pectolytic activity.

METHODS

Preparation of Oligogalacturonates and Analyses of Reaction Products

Oligogalacturonates with two to seven GalpA units were prepared from polygalacturonic acid (PGA), as described previously (Kester and Visser, 1990). Reduced oligogalacturonates were prepared according to the method of Omran et al. (1986). Quantitation of the oligomers and analyses of the reaction products were conducted as described previously (Parenicova et al., 1998). Enzymatic reaction rates for oligogalacturonates with different degrees of polymerization were determined spectrophotometrically at 235 nm by using 0.5 mM oligogalacturonate in 0.1 M 2-amino-2-methyl-1-propanol buffer, pH 9.5, in the presence of 1.0 mM CaCl₂ at 25°C. Enzyme activities were expressed as $\mu\text{kat}/\text{mg}$ by using the molar extinction coefficient at

235 nm for unsaturated digalacturonate of $4600 \text{ M}^{-1} \text{ cm}^{-1}$ at pH 8.0 (MacMillan and Vaughn, 1964). For the determination of bond cleavage frequencies, enzyme reactions were performed using the same reaction conditions, except that the buffer strength was lowered to 20 mM to prevent buffer component interference in the chromatographic analysis. Aliquots were taken at timed intervals, and the reactions were stopped by lowering the pH to 4.5 by the addition of 0.1 volume 1% acetic acid. Reaction products were analyzed as described previously (Hotchkiss et al., 1991; Hotchkiss and Hicks, 1993; Lieker et al., 1993; Benen et al., 1996) by using a Dionex BioLC high-performance chromatography system (Sunnyvale, CA). Detection was done by pulsed amperometry and spectrophotometry at 235 nm. Quantitation of the saturated reaction products was done using the amperometric data and the amperometric response of a calibration mixture of oligogalacturonates with different degrees of polymerization. The unsaturated oligogalacturonates were quantitated using the spectrophotometric response.

A fast-atom bombardment mass spectrum of pentaGalpA displayed the major difference between the molecular ion and hydrogen, (M-H)⁻, at a mass-to-charge ratio (*m/z*) of 897.6. Relatively minor amounts of (M-H)⁻ ions at *m/z* ratios of 721.5, 545.4, and 369.3 also were detected, representing tetraGalpA, triGalpA, and diGalpA, respectively. Mass spectra were obtained with a ZAB 2-SE high-field magnetic sector mass spectrometer (VG Analytical,

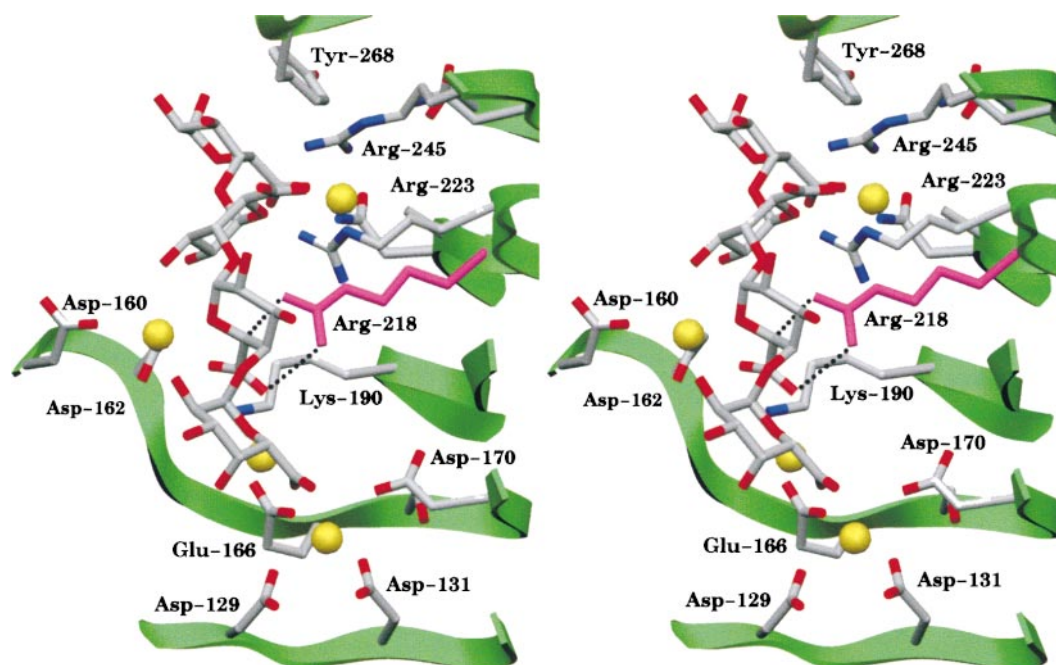


Figure 5. Stereoview of the $(\text{Ca}^{2+})_4$ -TetraGalpA Substrate Superimposed upon the Structure of Wild-Type PeIC.

The color scheme is the same as that used in Figure 1, except that wild-type Arg-218 is present and highlighted in magenta. One of the guanidinium nitrogens of Arg-218 lies within 2.6 Å of C-5 of GalpA³, and the other nitrogen lies within 2.7 Å of O-6B of GalpA³.

Manchester, UK) by using a cesium gun for fast-atom bombardment ionization at 8000 electron volts, glycerol-thioglycerol-triethylamine (10:10:1) as the matrix, and cesium iodide for mass calibration. The pentaGalpA sample was dissolved in water.

Preparation of Crystals

The R218K mutant of pectate lyase C (PeIC) was isolated from the periplasm of *Escherichia coli* HMS174(DE3) cells harboring pRSET5A constructs and purified as previously described (Kita et al., 1996). Crystals were grown using conditions similar to that for wild-type PeIC crystals (Yoder et al., 1990). The R218K mutant crystals are isomorphous with wild-type PeIC crystals and belong to space group P2₁2₁2₁ with unit cell parameters of $a = 72.14$ Å, $b = 78.32$ Å, and $c = 94.43$ Å, with one molecule per asymmetric unit. The R218K crystals were transferred from ammonium sulfate to a solution at pH 9.5 containing cryogenic agents, 7 mM Ca²⁺, and 50 mM pentaGalpA, and after 30 hr, they were frozen in liquid nitrogen for data collection.

X-Ray Diffraction Data Collection

X-ray diffraction data were collected to a resolution of 2.19 Å at -170°C by using a wavelength of 1.08 Å on a MARS imaging plate detector on Beam-Line 7-1 at the Stanford Synchrotron Radiation Light Source (Stanford, CA). The data were processed using MOS-FLM (Leslie, 1996). Data are included in Table 5.

Structure Determination

The structure has been solved by difference Fourier methods. Structure factors, by using the refined PeIC model (Yoder and Journak, 1995) in which Arg-218 had been omitted, were used to calculate a simulated annealed OMIT electron density map. Before water molecules were positioned, Ca²⁺ ions were fitted into the three highest peaks and refined without restraints. Four GalpA units were fitted to the remaining clustered density with the program O (Jones et al., 1991). The model was refined using the method of slow-cooling simulated annealing as implemented by X-PLOR (Brünger, 1996). The reflection data, with structure factor amplitudes (F) greater than two standard deviations, were randomly divided into two sets. The working set was composed of 90% of the data sampled at random, and the test set was composed of the remaining 10% of the data used for cross-validation of the refinement cycles (Brünger, 1993). The parameter and topology files used in X-PLOR were those of Engh and Huber (1991) for the protein and those of Ha et al. (1988), as modified by Weis et al. (1990), for the saccharide.

The conformation of each amino acid in the substrate binding region was adjusted by a series of refined OMIT maps in which a region of 8 Å around a residue had been omitted from the structure factor calculations and refinement (Hodel et al., 1992). In subsequent difference maps, peaks $>4\sigma$ and satisfying reasonable distance and geometry criteria were assigned as water molecules by using MAPMAN (Kleywegt and Jones, 1996). All water molecules were inspected visually. In one location, a water molecule could not account adequately for the residual electron density. Because the density was

Table 5. Crystallographic Data Collection and Refinement Statistics for the PelC R218K-(Ca²⁺)₃₋₄-PentaGalpA Complexes

Parameter	Value
Resolution	2.2 Å
Total observations	93,630
Unique observations	27,518
Percent completeness	95.1%
Average I/σ	11.8
R _{sym} ^a	3.1%
Resolution range of refinement	2.2 to 10.0 Å
No. of reflections with F > 2σ	26,662
Nonhydrogen protein atoms/asymmetric unit	2,647
Water molecules/asymmetric unit	328
Ca ²⁺ molecules/asymmetric unit ^b	3 to 4
GalpA molecules/asymmetric unit	4
R _{test} ^c for 10% data	23.3
R _{work} ^d for 90% data	18.6
Root-mean-square deviation from ideal geometry	
Bond length	0.0006 Å
Bond angle	1.54°
Impropers	1.17°
Average thermal factors	
Main chain	9.6 Å ²
Side chain	10.0 Å ²
All protein atoms	9.7 Å ²
Water molecules	22.6 Å ²
All nonhydrogen atoms	11.1 Å ²

^a R_{sym} = 100 = Σ|I_{avg} - I_{obs}|/ΣI_{avg}, when I_{avg} is the average (avg) or the observed (obs) intensity of the reflection.

^b Three of the Ca²⁺ ions refined to a 100% occupancy, and the fourth Ca²⁺ ion refined to a 50% occupancy.

^c R_{test} = Σ|F_o - F_c|/Σ|F_c| for the 10% of the reflections that were set aside for cross-validation and not used in the refinement.

^d R_{work} = Σ|F_o - F_c|/Σ|F_c| for the 90% of the reflections used in the refinement calculations.

located near to the carboxyl group of GalpA¹, in the same relative location as the other Ca²⁺ ions to the coordinating GalpA units, the density was assigned as a fourth Ca²⁺ ion, ¹Ca²⁺, with a lowered occupancy. After several cycles of refinements, the best fit of the density appeared to be that for a Ca²⁺ ion with 50% occupancy and a thermal factor of 15 Å². However, with this assignment, there remains an unfavorable contact with a well-ordered and fully occupied lysine, Lys-218. It is not possible to determine whether ¹Ca²⁺ interacts with an unprotonated lysine at pH 9.5 in the crystals in 50% of the molecules.

The final atomic model consists of 352 amino acids, 328 water molecules, 3.5 Ca²⁺ ions, and an α-1,4-linked oligosaccharide consisting of four well-ordered GalpA units. The final model refined to a crystallographic R factor of 18.6% for all measured reflections with structure-factor amplitudes F > 2σ in the 2.2 to 10.0 Å range. For the final statistics calculations, all reflections >2σ were used, including the test data set. The model was checked throughout using PROCHECK (Laskowski et al., 1993), and with the exception of the Lys-218-¹Ca²⁺ distance, no bad contacts were observed in the final model. Refinement statistics are summarized in Table 5. All model figures were prepared using SETOR (Evans, 1993).

Estimation of pK_a Values

The pK_a values for individual amino acids within the three-dimensional structure of the R218K-(Ca²⁺)₃₋₄-GalpA₅ complexes were determined using macroscopic electrostatics with atomic detail (MEAD), version 1.1.8 (Bashford and Gerwert, 1992). Standard partial atom charges were used, and the intrinsic pK_a values were calculated from the MEAD program multiflex.

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Structure of a Plant Cell Wall Fragment Complexed to Pectate Lyase C

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