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# Unique Substrate Recognition Mechanism of the Botulinum Neurotoxin D Light Chain\*

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**Background:** The mechanism of botulinum neurotoxin D light chain (LC/D) substrate recognition is not well defined.

**Results:** A dual recognition strategy employed by LC/D was revealed, in which one site of VAMP-2 was recognized by two independent, functionally similar LC/D sites that were complementary to each other.

**Conclusion:** LC/D utilizes a unique mechanism for substrate recognition.

**Significance:** This study provides insights for LC/D engineering and antitoxin development.

Botulinum neurotoxins are the most potent protein toxins in nature. Despite the potential to block neurotransmitter release at the neuromuscular junction and cause human botulism, they are widely used in protein therapies. Among the seven botulinum neurotoxin serotypes, mechanisms of substrate recognition and specificity are known to a certain extent in the A, B, E, and F light chains, but not in the D light chain (LC/D). In this study, we addressed the unique substrate recognition mechanism of LC/D and showed that this serotype underwent hydrophobic interactions with VAMP-2 at its V1 motif. The LC/D B3, B4, and B5 binding sites specifically recognize the hydrophobic residues in the V1 motif of VAMP-2. Interestingly, we identified a novel dual recognition mechanism employed by LC/D in recognition of VAMP-2 sites at both the active site and distal binding sites, in which one site of VAMP-2 was recognized by two independent, but functionally similar LC/D sites that were complementary to each other. The dual recognition strategy increases the tolerance of LC/D to mutations and renders it a good candidate for engineering to improve its therapeutic properties. In conclusion, in this study, we identified a unique multistep substrate recognition mechanism by LC/D and provide insights for LC/D engineering and antitoxin development.

The seven existing botulinum neurotoxin (BoNT)<sup>2</sup> serotypes, A–G, are the most potent protein toxins. BoNT causes human and animal botulism, a flaccid paralysis caused by the blocking of neurotransmitter (usually acetylcholine) release at the neuromuscular junction. Among the seven different serotypes, BoNT/A, BoNT/B, BoNT/E, and BoNT/F are involved in human botulism, whereas BoNT/D is responsible mainly for animal botulism (1–3).

BoNTs are zinc-dependent proteases and contain a His-Glu-X-X-His zinc-binding motif of metalloendopeptidases in the central region of their light chains (4). BoNTs are 150-kDa dichain proteins with typical A (active)-B (binding) structure-

function. BoNTs are activated by proteolysis to generate dichain organization (2). There are three domains found for BoNTs, including an N-terminal catalytic domain (light chain (LC)), an internal translocation domain (heavy chain), and a C-terminal receptor-binding domain (heavy chain) (5). The SNARE complex, formed by the vesicle SNARE (VAMP-2 (vesicle-associated membrane protein-2)) and the plasma membrane SNAREs (SNAP25 and syntaxin 1a), is the driving force of vesicle fusion during the mammalian neuronal exocytosis process (6). These SNARE proteins are the targets of the seven BoNTs: serotypes B, D, F, and G cleave VAMP-2; serotypes A and E cleave SNAP25; and serotype C cleaves both SNAP25 and syntaxin 1a (2).

Due to their extreme toxicity and ease of production, handling, and delivery through aerosol or liquid route, BoNTs represent a potential biological warfare agent and have been classified as a category A agent by the Center for Disease Control and Prevention in the United States (7, 8). However, another key feature of BoNTs is that their intoxication can be reversed by the replacement of affected nerves with new ones (5, 9, 10), thus making BoNTs effective agents for the therapies of a range of neuromuscular disorders such as strabismus (11, 12). A better understanding of the mechanism of action and substrate recognition of BoNTs will enable us to develop an antidote for BoNT intoxication and novel therapies to extend its clinical applications. Recently, the mechanisms of BoNT/A, BoNT/B, BoNT/E, and BoNT/F have been thoroughly studied, with results indicating that an extended region (exosite) is necessary for substrate binding and cleavage (13–16). In 2006, Arndt *et al.* (17) reported the structure of the light chain of BoNT/D and proposed the recognition of the hydrophobic SNARE V1 motif through structural comparison with LC/F. Sikorra *et al.* (18) reported the contribution of VAMP-2 residues to LC/D substrate recognition and found that LC/D requires a relatively short sequence for optimal substrate cleavage compared with other serotypes such as LC/F and the tetanus neurotoxin LC. However, the mechanism underlying the effective recognition and cleavage of VAMP-2 by LC/D is still unclear, which prompted us in this study to depict the mechanism of substrate recognition and specificity by LC/D.

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<sup>2</sup> The abbreviations used are: BoNT, botulinum neurotoxin; LC, light chain.

## VAMP-2 Recognition by BoNT/D

Our data show that, similar to other serotypes, LC/D recognition of VAMP-2 occurs through multistep binding and, in particular, recognition of VAMP-2 sites by LC/D substrate recognition pockets. Interestingly, in contrast to all other BoNT LCs and metalloproteinases, LC/D employs a novel dual recognition mechanism, in which one VAMP-2 site is recognized by two independent LC/D sites that are complementary to each other. The dual recognition strategy increases the tolerance of LC/D to mutations and makes it a good candidate for engineering to improve its pharmacological properties.

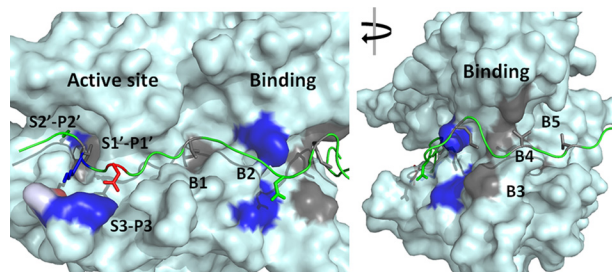
### EXPERIMENTAL PROCEDURES

**Molecular Modeling**—The structure of the LC/D·VAMP-2 complex was modeled and analyzed using SWISS-MODEL and refined using PyMOL as described previously (15).

**Plasmid Construction and Protein Expression**—Codon-optimized DNA encoding the LC domain of BoNT/D (residues 1–430) was synthesized by EZBiolab (Westfield, IN). The PCR product was then subcloned into the pET-15b vector and transformed into *Escherichia coli* BL21(DE3)-RIL cells (Stratagene). Protein expression and purification were achieved as described previously (14, 15, 19). VAMP-2(1–97) was constructed by PCR amplification of the cDNA clone purchased from American Type Culture Collection (GenBank™ accession number NM\_014232) and subcloning into pGEX-2T. The plasmid containing the VAMP-2(1–97) fragment was transformed into *E. coli* BL21(DE3) cells, and expression and purification of VAMP-2 were achieved as described previously (14, 15, 19).

**VAMP-2 and LC/D Mutagenesis**—The introduction of point mutations into LC/D and VAMP-2 genes was also performed using the QuikChange® protocol as described previously (15, 16). Plasmids were sequenced to confirm the mutations and that additional mutations were not present within the open reading frame of VAMP-2 and LC/D. Mutant proteins were produced and purified as described previously (15, 16).

**Linear Velocity and Kinetic Constants**—Determination of the linear velocity and kinetic constants of LC/D and its derivatives was performed as described previously (14, 15, 19). Briefly, 10  $\mu\text{M}$  VAMP-2 or the indicated VAMP-2 derivatives were incubated with various concentrations of LC/D or its derivatives in 10  $\mu\text{l}$  of reaction buffer (10 mM Tris-HCl (pH 7.6) and 20 mM NaCl) at 37 °C for 20 min. The reactions were stopped by adding an equal volume of SDS-PAGE sample buffer, boiled at 100 °C for 5 min, and analyzed for the relative abundance of the substrate and cleaved product by SDS-PAGE (12% polyacrylamide gels). The amount of cleaved VAMP-2 was determined by densitometry.  $K_m$  and  $k_{\text{cat}}$  determinations were performed using the same assay, in which VAMP-2 concentrations were adjusted to between 1 and 20  $\mu\text{M}$  to achieve ~10% cleavage by LC/D or its derivatives. The amount of cleaved VAMP-2 substrate was determined by densitometry, and the velocity was determined by dividing the amount of substrate cleavage by the reaction time. The reaction velocity against the substrate concentration was fitted to the Michaelis-Menten equation, and kinetic constants were derived using GraphPad. At least five independent assays were performed to determine the kinetic constants for each protein.



**FIGURE 1. Overall view of the modeled LC/D·VAMP-2 complex structure.** Left panel, view of the active site side; right panel, view after a 90° clockwise turn. LC/D is shown as a surface structure, and VAMP-2 is shown as a ribbon structure. The active site recognition and binding site interactions are highlighted. Negatively charged residues are shown in red, positively charged residues are shown in blue, hydrophobic residues are shown in gray, and polar residues are shown in green.

**Trypsin Digestion of LC/D and Its Derivatives**—10  $\mu\text{M}$  LC/D and its derivatives were incubated with 2 mM trypsin in a 20- $\mu\text{l}$  reaction volume at 37 °C for 30 min. The reactions were stopped by adding SDS-PAGE sample buffer, subjected to SDS-PAGE, and stained to visualize the partial trypsin digestion profiles.

**Far-UV Circular Dichroism Analysis**—LC/D and its derivatives were subjected to far-UV CD analysis. CD spectroscopy was performed at a wavelength range of 200–250 nm at room temperature with a JASCO J-810 spectropolarimeter. Far-UV CD data were obtained with a 10-mm path length quartz cuvette containing 500  $\mu\text{l}$  of protein solution (0.1–0.4 mg/ml protein in 10 mM Tris-HCl (pH 7.9) and 20 mM NaCl) at a scanning speed of 50 nm/min and a 2-s response time. Each sample was measured in triplicate, the CD data were converted to molar ellipticity, and the spectrum was generated using GraphPad.

### RESULTS

A previous study has shown that several residues of VAMP-2 contribute to substrate cleavage by LC/D, including Val<sup>39</sup>, Val<sup>42</sup>, Met<sup>46</sup>, Val<sup>49</sup>, Asp<sup>53</sup>, Lys<sup>59</sup>, Leu<sup>60</sup>, and Ser<sup>61</sup> (18). To better quantify the degree of contribution of these residues to LC/D substrate cleavage, linear velocity assays were performed to depict the impact of alanine mutagenesis of these residues. The results show that amino acid changes in VAMP-2, including V39A, V42A, M46A, V49A, D53A, D57A, K59A, L60A, and S61A, caused ~25-, 10-, 125-, 20-, 25-, 10-, 20-, 25-, and 20-fold reductions in LC/D cleavage, respectively. These data confirm that the residues in VAMP-2 contribute significantly to LC/D substrate recognition and cleavage. Analysis of the modeled structure of the LC/D·VAMP-2 complex identified three putative substrate recognition pockets in the active site of LC/D, S2', S1', and S3, which may specifically recognize the VAMP-2 P2' (Ser<sup>61</sup>), P1' (Leu<sup>60</sup>), and P3 (Asp<sup>53</sup>) sites, respectively (Fig. 1). In addition, we also predicted several other substrate-binding pockets distal to the active site of LC/D, including the B1–B5 binding sites (Fig. 1). To characterize the substrate recognition pockets and to confirm the specific recognition of VAMP-2 sites, the LC/D residues constituting the substrate recognition pockets were mutated to different amino acid residues to test the effects of these changes on substrate recognition. To exclude the possibility that the effect was due to the



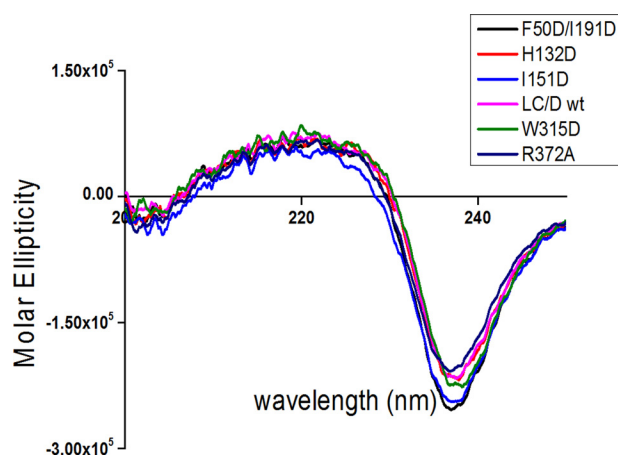


FIGURE 2. **CD spectroscopy analysis of LC/D and its derivatives.** Far-UV CD (200–250 nm) data were obtained for LC/D and its derivatives with a JASCO J-810 spectropolarimeter at room temperature. The data for the most representative LC/D derivatives are shown in different colors.

overall conformational changes as a result of the mutations, we performed partial trypsin digestion on different mutant proteins, and the result indicated that all of the LC/D mutants had an identical digestion profile to WT-LC/D, suggesting that the mutations did not cause any conformational change in LC/D (data not shown). In addition, far-UV CD analysis of LC/D and its derivatives indicated that LC/D(I151D) had a slightly different spectrum compared with WT-LC/D, whereas all other LC/D derivatives had the same far-UV CD spectrum as WT-LC/D (Fig. 2). The curve vertexes of LC/D and its derivatives at  $\sim 240$  nm looked different, which is probably due to the high degree of flexibility of LC/D owing to its relatively high number of turns and random coils. Therefore, the different curves at  $\sim 240$  nm did not reflect the conformational changes of LC/D derivatives (Fig. 2). The CD spectra of the other LC/D derivatives were similar to that of WT-LC/D (data not shown).

#### Recognition of the P2' Site (Ser<sup>61</sup>) of VAMP-2 by the S2' Pocket of LC/D

The P2' site (Ser<sup>61</sup>) of VAMP-2 plays a certain role in LC/D substrate recognition, as the VAMP-2(S61A) mutation reduced LC/D substrate hydrolysis by  $\sim 20$ -fold. A S2' pocket in LC/D that recognized the P2' site of VAMP-2 at Ser<sup>61</sup> was identified through analysis of the modeled structure of the complex of LC/D and VAMP-2. The S2' pocket is composed of Arg<sup>372</sup>, and the R372A mutation resulted in an  $\sim 40$ -fold reduction of LC/D activity, with almost the same  $K_m$  and an  $\sim 40$ -fold lower  $k_{cat}$  compared with WT-LC/D (Fig. 3a and Table 1). These data suggest that the S2' pocket (Arg<sup>372</sup>) of LC/D may recognize the P2' site of VAMP-2 by forming a hydrogen bond between these two residues.

#### Dual Recognition of the VAMP-2 P1' Site by the S1' Pocket of LC/D

The P1' site of VAMP-2 was shown to be important for LC/D substrate recognition, as seen in other serotypes of BoNT (14, 15, 20, 21). Mutation of the VAMP-2 P1' residue (L60A) reduced LC/D substrate hydrolysis by  $\sim 25$ -fold. A corresponding S1' pocket composed of two hydrophobic residues, Tyr<sup>168</sup>

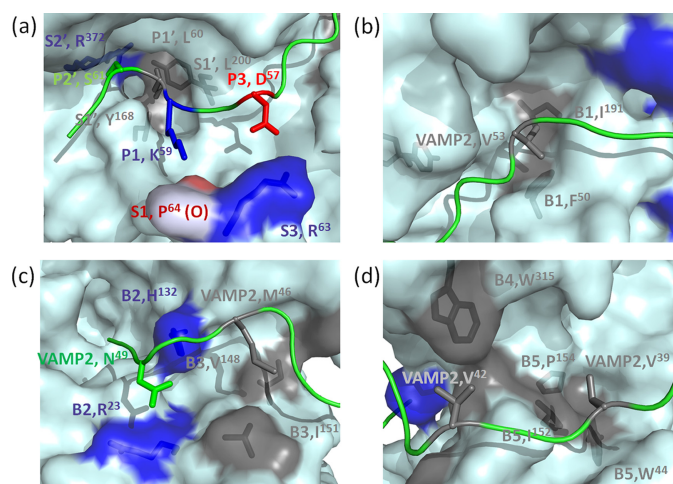


FIGURE 3. **Specific recognition of VAMP-2 by LC/D pockets.** Shown are surface representations of the recognition of different P sites of VAMP-2 by the S pockets and recognition of VAMP-2-binding sites by the B1–B5 binding sites of LC/D. Negatively charged residues are shown in blue, positively charged residues are shown in red, hydrophobic residues are shown in gray, and polar residues are shown in green. *a*, recognition of the P sites of VAMP-2 by the active site pockets of LC/D. The P2' site (Ser<sup>61</sup>) of VAMP-2 is recognized by the S2' pocket (Arg<sup>372</sup>). The P1' site (Leu<sup>60</sup>) of VAMP-2 is recognized by the S1' pocket (Tyr<sup>168</sup> and Leu<sup>200</sup>) of LC/D. The P1 site (Lys<sup>59</sup>) of VAMP-2 interacts with the oxygen atom of Pro<sup>64</sup> of LC/D. The P3 site (Asp<sup>57</sup>) of VAMP-2 is recognized by the S3 pocket (Arg<sup>63</sup>) of LC/D. *b*, recognition of Val<sup>53</sup> by the B1 binding site (Phe<sup>50</sup> and Ile<sup>191</sup>) of LC/D. *c*, recognition of Asn<sup>49</sup> of VAMP-2 by the B2 binding site (Arg<sup>23</sup> and His<sup>132</sup>) of LC/D and recognition of Met<sup>46</sup> of VAMP-2 by the B3 binding site (Val<sup>148</sup> and Ile<sup>151</sup>) of LC/D. *d*, recognition of Val<sup>42</sup> by the B4 binding site (Trp<sup>315</sup>) of LC/D and recognition of Val<sup>39</sup> by the B5 binding site (Trp<sup>44</sup>, Ile<sup>152</sup>, and Pro<sup>154</sup>) of LC/D.

and Leu<sup>200</sup>, was identified in LC/D (Fig. 3a). The LC/D(Y168A) mutation had no effect on LC/D substrate hydrolysis, and LC/D(Y168D) affected substrate hydrolysis by only  $\sim 2$ -fold. The LC/D(L200A) mutation resulted in an  $\sim 2$ -fold reduction of LC/D substrate hydrolysis, whereas LC/D(L200D) affected substrate hydrolysis by  $\sim 8$ -fold (Table 1). The complementary effect of Tyr<sup>168</sup> and Leu<sup>200</sup> was also examined, and we found that although LC/D(Y168A/L200A) resulted in only an  $\sim 2$ -fold reduction of substrate hydrolysis, LC/D(Y168D/L200D) reduced substrate hydrolysis by 60-fold, with no effect on  $K_m$  and an  $\sim 60$ -fold reduction of  $k_{cat}$ . These data suggest that a hydrophobic S1' pocket is necessary to maintain the full recognition of VAMP-2 Leu<sup>60</sup> and that both hydrophobic residues in the S1' pocket, Tyr<sup>168</sup> and Leu<sup>200</sup>, play a complementary role in Leu<sup>60</sup> recognition.

#### The S3 Pocket Residue of LC/D Interacts with the P3 Residue (Asp<sup>57</sup>) of VAMP-2

Asp<sup>57</sup> at the P3 site of VAMP-2 plays a certain role in LC/D substrate recognition, as the D57A mutation reduced LC/D substrate hydrolysis by  $\sim 20$ -fold. The S3 pocket of LC/D that specifically recognized the P3 site residue (Asp<sup>57</sup>) of VAMP-2 contains Arg<sup>63</sup>. The R63A mutation had almost no effect on  $K_m$ , but reduced substrate catalysis by  $\sim 13$ -fold (Fig. 3a and Table 1). The charge reversal mutation LC/D(R63E) caused an  $\sim 50$ -fold reduction of LC/D substrate hydrolysis, with no effect on  $K_m$  and an  $\sim 50$ -fold reduction of  $k_{cat}$ . These data suggest that a salt bridge or a side chain hydrogen bond is important for recognition of Asp<sup>57</sup> of VAMP-2 by Arg<sup>63</sup> of LC/D.

**TABLE 1**  
Efficiency of VAMP-2 hydrolysis and kinetic constants of LC/D and its derivatives

LC/D pockets	VAMP-2 site recognition	LC/D derivatives	Activity reduction <sup>a</sup>	$K_m$	$k_{cat}$	$k_{cat}/K_m$
		WT- LC/D	1	$\mu M$	$s^{-1}$	$s^{-1} \mu M^{-1}$
			<i>-fold</i>			
AS-S2 <sup>c</sup>	P2' (Ser <sup>61</sup> )	R372A	40	2.91 (0.74) <sup>b</sup>	6.88	2.36
AS-S1'	P1' (Leu <sup>60</sup> )	Y168A	1	— <sup>d</sup>	—	—
		Y168D	2	—	—	—
		L200A	2	—	—	—
		L200D	8	2.40 (0.36)	0.81	$3.4 \times 10^{-1}$
		Y168A/L200A	2	—	—	—
		Y168D/L200D	60	3.33 (0.66)	0.12	$3.6 \times 10^{-2}$
AS-S3	P3 (Asp <sup>57</sup> )	R63A	10	2.19 (0.24)	0.40	$1.8 \times 10^{-1}$
		R63E	50	2.26 (0.35)	0.15	$6.6 \times 10^{-2}$
Binding B1	Val <sup>53</sup>	F50A	1	—	—	—
		I191A	1	—	—	—
		F50D	4	3.24 (0.12)	1.60	$4.9 \times 10^{-1}$
		I191D	4	3.56 (0.21)	1.40	$3.9 \times 10^{-1}$
		F50A/I191A	60	4.81 (0.64)	0.28	$5.8 \times 10^{-2}$
		F50D/I191D	400	4.46 (0.57)	0.04	$8.9 \times 10^{-3}$
Binding B2	Asn <sup>49</sup>	R23A	1	—	—	—
		H132A	1	—	—	—
		H132D	ND <sup>d</sup>	ND <sup>b</sup>	ND	ND
		H132Q	100	35.56 (3.48)	0.86	$2.4 \times 10^{-2}$
		R23D/H132A	25	32.87 (5.25)	2.38	$7.2 \times 10^{-2}$
Binding B3	Met <sup>46</sup>	V148A	1	—	—	—
		V148D	1	—	—	—
		I151A	2	—	—	—
		I151D	1000	7.96 (1.284)	0.02	$2.5 \times 10^{-3}$
		V148/I151A	15	—	—	—
Binding B4	Val <sup>42</sup>	W315A	20	9.90 (1.51)	1.52	$1.5 \times 10^{-1}$
		W315D	40	47.63 (15.12)	3.92	$8.2 \times 10^{-2}$
Binding B5	Val <sup>39</sup>	W44D	2	—	—	—
		I152D	2	—	—	—
		P154D	4	11.86 (3.46)	6.35	$5.4 \times 10^{-1}$
		W44A/I152A/P154A	20	32.45 (8.87)	5.86	$1.8 \times 10^{-1}$

<sup>a</sup> WT-VAMP-2 hydrolysis was measured as the ratio of the amount of LC/D derivatives needed to cleave 50% of WT-VAMP-2 to the amount of WT-LC/D needed to cleave 50% of WT-VAMP-2.

<sup>b</sup> The numbers in parentheses are the S.E. of at least five independent experiments.

<sup>c</sup> AS, active site; ND, not detectable. The mutant was too inactive to determine its kinetic constants in our experiments.

<sup>d</sup> —, kinetic constants were not determined.

### The Main Chain Oxygen Atom of LC/D Pro<sup>64</sup> Interacts with the VAMP-2 P1 Site Residue (Lys<sup>59</sup>)

There was no obvious residue or pocket that showed an interaction with the P1 site residue (Lys<sup>59</sup>) of VAMP-2. The VAMP-2(K59A) mutation caused an ~20-fold reduction of LC/D substrate hydrolysis, suggesting a role for Lys<sup>59</sup> in LC/D substrate recognition. Structural analysis indicated that Lys<sup>59</sup> could potentially interact with the oxygen atom of LC/D Pro<sup>64</sup> through formation of a hydrogen bond. This interaction could not be tested through mutational analysis (Fig. 3a).

### Substrate Binding and Recognition Distal to the Active Site of LC/D

**Recognition of VAMP-2 Val<sup>53</sup> by the LC/D B1 Binding Site**—The VAMP-2(V53A) mutation caused an ~25-fold reduction of LC/D hydrolysis. The B1 binding site of LC/D, which is formed by two residues, Phe<sup>50</sup> and Ile<sup>191</sup>, may interact with Val<sup>53</sup> (Fig. 3b). The LC/D(F50A) and LC/D(I191A) mutations had no effect on LC/D substrate hydrolysis, whereas LC/D(F50D) and LC/D(I191D) caused an ~4-fold reduction of VAMP-2 hydrolysis. To test for the complementary effect of these two residues on Val<sup>53</sup> recognition, the effects of double

mutations were tested. The double mutation LC/D(F50A/I191A) resulted in an ~60-fold reduction of substrate hydrolysis, with an ~2-fold increase in  $K_m$  and an ~25-fold decrease in  $k_{cat}$ , whereas LC/D(F50D/I191D) reduced substrate hydrolysis by ~400-fold, with an ~2-fold increase in  $K_m$  and an ~200-fold decrease in  $k_{cat}$  (Table 1). These data suggest that Phe<sup>50</sup> and Ile<sup>191</sup> have independent but complementary effects on Val<sup>53</sup> substrate recognition. The recognition of Val<sup>53</sup> might contribute mainly to the fine orientation of VAMP-2 for optimal substrate recognition in the active site of LC/D because this recognition site contributed mainly to substrate catalysis ( $k_{cat}$ ), but not substrate binding ( $K_m$ ).

**Recognition of VAMP-2 Asn<sup>49</sup> by the LC/D B2 Binding Site**—The VAMP-2(N49A) mutation was associated with an ~20-fold reduction in cleavage efficiency of LC/D. The B2 binding site, which is composed of Arg<sup>23</sup> and His<sup>132</sup> of LC/D and may interact with Asn<sup>49</sup>, was revealed through analysis of the structure of the LC/D-VAMP-2 complex (Fig. 3c). The LC/D(R23A) and LC/D(H132A) mutations had no effect on LC/D substrate hydrolysis (Table 1). Surprisingly, the charge reversal mutation LC/D(R23D) still maintained the full activity on VAMP-2 as did WT-LC/D. The charge reversal mutation LC/D(H132D)

became inactive in cleaving VAMP-2, whereas LC/D(H132Q) reduced substrate hydrolysis by  $\sim 100$ -fold (Table 1), suggesting that the formation of a hydrogen bond between LC/D His<sup>132</sup> and VAMP-2 Asn<sup>49</sup> may contribute to this recognition and that a negatively charged residue at position 132 may impair this interaction. However, the minimal effect of the LC/D(H132A) mutation on LC/D substrate hydrolysis may be due to the complementary effect of Arg<sup>23</sup>. To test this hypothesis, the effect of R23D and H132A double mutations on LC/D substrate hydrolysis was tested. LC/D(R23D/H132A) reduced  $K_m$  by  $\sim 11$ -fold and  $k_{cat}$  by  $\sim 3$ -fold (Table 1), suggesting that both Arg<sup>23</sup> and His<sup>132</sup> play a role in LC/D substrate catalysis and that His<sup>132</sup> plays a dominant role in Asn<sup>49</sup> recognition. This substrate recognition contributes significant to substrate binding.

**Recognition of VAMP-2 Met<sup>46</sup> by the LC/D B3 Binding Site**—Compared with Asn<sup>49</sup>, Met<sup>46</sup> of VAMP-2 plays a more important role in VAMP-2 recognition by LC/D. Consistent with previous data (18), under our assay conditions, the VAMP-2(M46A) mutation reduced LC/D substrate hydrolysis by  $\sim 125$ -fold. Structural analysis also identified the B3 binding site, which is composed of two residues, Ile<sup>151</sup> and Val<sup>148</sup>, and which may interact with Met<sup>46</sup> (Fig. 3c). The point mutations associated with the LC/D(V148A) and LC/D(I151A) alterations did not show any effect on VAMP-2 hydrolysis. LC/D(V148D) also did not have any impact on VAMP-2 hydrolysis, but LC/D(I151D) reduced LC/D substrate hydrolysis by  $\sim 1000$ -fold (Table 1). The dramatic effect of the LC/D(I151D) mutation may be partially due to its conformational change based on our far-UV CD analysis. Similar to Asn<sup>49</sup> recognition, the minimal effect of the LC/D(I151A) mutation may be related to the complementary effect of Val<sup>148</sup>. To test this hypothesis, the effect of Ile<sup>151</sup> and Val<sup>148</sup> double mutations on LC/D substrate hydrolysis was tested. LC/D(I151A/V148D) reduced  $K_m$  by  $\sim 10$ -fold and  $k_{cat}$  by  $\sim 2$ -fold (Table 1), suggesting that both Ile<sup>151</sup> and Val<sup>148</sup> play a role in LC/D substrate catalysis, with Ile<sup>151</sup> playing a dominant role in substrate recognition. This substrate recognition contributes significantly to substrate binding.

**Recognition of VAMP-2 Val<sup>42</sup> by the LC/D B4 Binding Site**—Val<sup>42</sup> of VAMP-2 is also important for LC/D substrate hydrolysis, and the VAMP-2(V42A) mutation affected LC/D cleavage of VAMP-2 by  $\sim 10$ -fold. Based on the modeled structure of the LC/D-VAMP-2 complex, LC/D Trp<sup>315</sup> in the B4 binding site was predicted to have a direct interaction with Val<sup>42</sup> (Fig. 3d). The LC/D(W315A) mutation reduced VAMP-2 hydrolysis by  $\sim 20$ -fold, with an  $\sim 4$ -fold increase in  $K_m$  and an  $\sim 5$ -fold decrease in  $k_{cat}$ , whereas LC/D(W315D) reduced VAMP-2 hydrolysis by  $\sim 40$ -fold, with an  $\sim 20$ -fold increase in  $K_m$  and an  $\sim 2$ -fold decrease in  $k_{cat}$ . These data suggest that LC/D Trp<sup>315</sup> and VAMP-2 Val<sup>42</sup> substrate recognition contributes significantly to substrate binding.

**Recognition of VAMP-2 Val<sup>39</sup> by the LC/D B5 Binding Site**—Another hydrophobic residue in VAMP-2 that contributes to LC/D substrate hydrolysis is Val<sup>39</sup>. The mutation VAMP-2(V39A) resulted in an  $\sim 25$ -fold reduction of LC/D hydrolysis. A hydrophobic pocket in LC/D, the B5 binding site, which is formed by Trp<sup>44</sup>, Ile<sup>152</sup>, and Pro<sup>154</sup>, was identified through analysis of the structure of the LC/D-VAMP-2 com-

plex (Fig. 3d). The W44D, P152D, and P154D mutations were associated with  $\sim 2$ -,  $2$ -, and  $4$ -fold reductions of substrate hydrolysis, respectively (Table 1). These mutations had no effect on  $k_{cat}$ , but caused a  $4$ -fold increase in  $K_m$ , suggesting a role for this pocket in substrate binding. Interestingly, the triple mutation LC/D(W44A/I152A/P154A) resulted in an  $\sim 20$ -fold increase in  $K_m$ , but had no effect on  $k_{cat}$  (Table 1), suggesting a complementary effect of these three residues on Val<sup>39</sup> recognition and VAMP-2 binding.

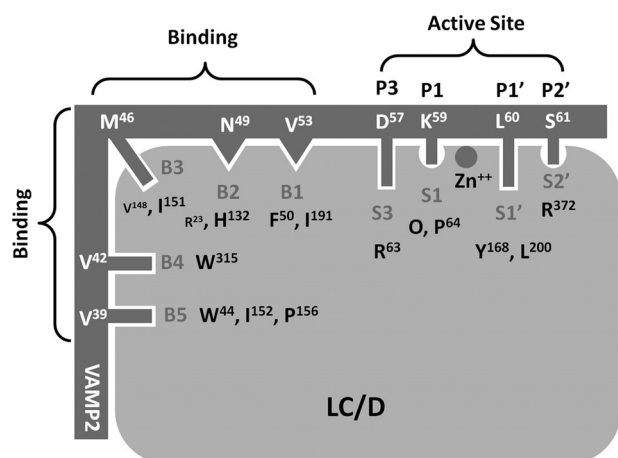
## DISCUSSION

It was proposed that botulinum neurotoxins recognize their substrates through two separate regions, one that contains the scissile bond and the other distal to the scissile bond and containing the SNARE motif (22). A two-region substrate recognition model has been demonstrated in LC/A, LC/B, LC/E, LC/F, and the tetanus neurotoxin LC (14, 15, 20, 21). The significance of the SNARE motif was not consistently proven in these toxins. However, for LC/D, the SNARE V1 motif (<sup>38</sup>QVDEVVDIMR<sup>47</sup>) was shown to be important for substrate recognition and hydrolysis. Three conservative hydrophobic residues in the V1 motif of VAMP-2, Val<sup>39</sup>, Val<sup>42</sup>, and Met<sup>47</sup>, are critical for efficient LC/D substrate hydrolysis. In addition, LC/D also utilizes other hydrophobic interactions to recognize the VAMP-2 substrate, such as the recognition of the P1' site (Leu<sup>60</sup>) of VAMP-2 by the hydrophobic S1' pocket of LC/D and the recognition of Val<sup>53</sup> of VAMP-2 by the hydrophobic B1 binding site of LC/D. In contrast to the substrate binding contributed by the SNARE V1 motif (B3, B4, and B5), the B1 and B2 binding sites contribute more to LC/D substrate catalysis than substrate binding. The data suggest that the V1 motif plays a significant role in LC/D substrate binding, whereas the B1 and B2 binding sites may help more in fine-tuning the orientation of the substrate for specific recognition by the active site of LC/D rather than direct substrate binding.

In this study, we revealed the mechanism of LC/D substrate recognition and specificity. After internalization to the cytoplasm of neuronal cells, LC/D attacks the free form of VAMP-2 through interaction with and recognition of hydrophobic residues in the V1 motif of VAMP-2, including Val<sup>39</sup>, Val<sup>42</sup>, and Met<sup>46</sup>, by the substrate-binding regions B5, B4, and B3 of LC/D on the substrate-binding cleft, respectively. In particular, binding of Met<sup>46</sup> of VAMP-2 to the LC/D B3 binding site was suggested to be very important for LC/D substrate recognition. This binding facilitates further binding of VAMP-2 Asn<sup>49</sup> and Val<sup>53</sup> to the B2 and B1 binding sites located at the active site surface of LC/D. The recognition of VAMP-2 Val<sup>53</sup> by the LC/D Phe<sup>50</sup>/Ile<sup>191</sup> pocket further orientates and stabilizes VAMP-2 for subsequent recognition of its different P sites by the corresponding S pockets in the active site of LC/D. Active P site recognition includes the formation of a salt bridge between P3 (Asp<sup>57</sup>) of VAMP-2 and S3 (Arg<sup>63</sup>) of LC/D, a hydrogen bond interaction between P1 (Lys<sup>58</sup>) of VAMP-2 and the main chain oxygen atom of Pro<sup>64</sup>, recognition of P1' (Leu<sup>60</sup>) of VAMP-2 by the S1' pocket (Tyr<sup>168</sup> and Leu<sup>200</sup>) of LC/D, and finally a hydrogen bond interaction between P2' (Ser<sup>61</sup>) of VAMP-2 and the S2' pocket (Arg<sup>372</sup>) of LC/D. The anchoring of VAMP-2 P sites to different S pockets in the active site of



## VAMP-2 Recognition by BoNT/D



**FIGURE 4. Mechanism of substrate recognition by LC/D.** After internalization to the cytoplasm of neuronal cells, LC/D attacks the free form of VAMP-2 through interaction with and recognition of hydrophobic residues in the V1 motif of VAMP-2, including Val<sup>39</sup>, Val<sup>42</sup>, and Met<sup>46</sup>, by the substrate-binding regions B5, B4, and B3 of LC/D on the substrate-binding cleft, respectively. In particular, binding of Met<sup>46</sup> of VAMP-2 to the LC/D B3 binding site was suggested to be very important for LC/D substrate recognition. This binding facilitates further binding of VAMP-2 Asn<sup>49</sup> and Val<sup>53</sup> to the B2 and B1 binding sites located at the active site surface of LC/D. The recognition of VAMP-2 Val<sup>53</sup> by the LC/D Phe<sup>50</sup>/Ile<sup>191</sup> pocket further orientates and stabilizes VAMP-2 for subsequent recognition of different P sites of VAMP-2 by the corresponding S pockets in the active site of LC/D. Active P site recognition includes the formation of a salt bridge between P3 (Asp<sup>57</sup>) of VAMP-2 and S3 (Arg<sup>63</sup>) of LC/D, a hydrogen bond interaction between P1 (Lys<sup>59</sup>) of VAMP-2 and the main chain oxygen atom of Pro<sup>64</sup>, recognition of P1' (Leu<sup>60</sup>) of VAMP-2 by the S1' pocket (Tyr<sup>168</sup> and Leu<sup>200</sup>) of LC/D, and finally a hydrogen bond interaction between P2' (Ser<sup>61</sup>) of VAMP-2 and the S2' pocket (Arg<sup>372</sup>) of LC/D. The anchoring of VAMP-2 P sites to different S pockets in the active site of LC/D aligns the VAMP-2 scissile bond close enough to the active site zinc ion to facilitate peptide bond cleavage.

LC/D aligns the VAMP-2 scissile bond close enough to the active site zinc ion to facilitate peptide bond cleavage (Fig. 4).

Compared with substrate recognition by other serotypes of BoNT, LC/D possesses unique features of substrate recognition (14, 15, 20, 21). First, hydrophobic interaction between LC/D and VAMP-2 plays an important role in substrate recognition. The interaction between VAMP-2 Met<sup>46</sup> and LC/D Ile<sup>151</sup> seems to be critical for LC/D substrate recognition. This may be the first step in substrate recognition, which may facilitate the conformational change in VAMP-2 from a double helix to a free loop confirmation, favoring the subsequent substrate binding and catalysis by different regions of LC/D. Further research may be needed to test this hypothesis. However, far-UV CD analysis showed that I151D displays a slightly different conformation compared with WT-LC/D, suggesting that the significant effect of I151D substrate recognition may be partially due to the conformational change in the whole protein, but not the loss of the Ile<sup>151</sup> site recognition. Second, in contrast to the recognition of one site of the substrate by one pocket of the LC for other serotypes of BoNT, LC/D utilizes two functionally similar residues to recognize one site of VAMP-2, such as the S1' pocket (Tyr<sup>168</sup>–Leu<sup>200</sup>) of LC/D for recognition of the P1' site (Leu<sup>60</sup>) of VAMP-2 in the active site of LC/D. In addition, dual recognition is also commonly employed at the substrate-binding regions. VAMP-2 Val<sup>53</sup> is recognized by the LC/D pocket formed by Phe<sup>50</sup> and Ile<sup>191</sup>. Mutation to each residue did not have much effect (maximum of 4-fold) on substrate hydroly-

ysis, whereas mutation to both residues resulted in a dramatic reduction (400-fold) of substrate hydrolysis. VAMP-2 Asn<sup>49</sup> and Met<sup>46</sup> are also recognized by pockets with the dual recognition mechanism. The pocket that recognizes Asn<sup>49</sup> of VAMP-2 is formed by Arg<sup>23</sup> and His<sup>132</sup> of LC/D. Although His<sup>132</sup> plays a dominant role in Asn<sup>49</sup> recognition, the H132A mutation can be complemented by Arg<sup>23</sup>. Similar to Asn<sup>49</sup>, the pocket that recognizes Met<sup>46</sup> of VAMP-2 is formed by Val<sup>148</sup> and Ile<sup>151</sup> of LC/D. Ile<sup>151</sup> plays a dominant role in Met<sup>46</sup> recognition, whereas Val<sup>148</sup> of VAMP-2 can play a complementary role when Ile<sup>151</sup> is mutated to alanine. Finally, the pocket that recognizes Val<sup>39</sup> of VAMP-2 is formed by three hydrophobic residues (Trp<sup>44</sup>, Ile<sup>152</sup>, and Pro<sup>154</sup>) of LC/D. Mutation of each residue to alanine or asparagine had no effect or only a minor effect on substrate hydrolysis, whereas triple mutations to alanine resulted in a much stronger reduction of substrate hydrolysis, highlighting the complementary effects of the three residues forming this pocket. The presence of two or more functionally similar residues in the same substrate recognition pocket enables LC/D to tolerate mutations. This property of LC/D makes it a good candidate for further protein engineering.

Unlike BoNT/A, which is the most toxic botulinum neurotoxin and is implicated in human botulism, BoNT/D is responsible mainly for animal botulism, such as cattle botulism. However, our data indicate that LC/D and LC/A exhibit a similar potency in hydrolyzing their substrates under *in vitro* conditions (16). The role of BoNT/D as a human therapy or bioterrorism weapon remains to be investigated. Our data provide insights into the development of novel BoNT-based therapies and BoNT/D antitoxins.

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