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	Vibrio cholerae Hemagglutinin/Protease Nicks Cholera Enterotoxin
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,%- <u></u>	Departments of Microbiology ¹ and Biochemistry, ² School of Medicine, University of Missouri, Columbia, Missouri 65212
	Received 27 March 1984/Accepted 22 May 1984
<u> </u>	Unnicked cholera enterotoxin was isolated from culture supernatants of Vibrio cholerae 569B by either rapid
	processing of flask-grown cultures or by growing and processing fermentor cultures in the presence of ethylene
	Processing of hask-grown cultures of by growing and drocessing termentor cultures in the dresence of ethylene
	glycol-bis(β -aminoethyl ether)- N , N' , N' -tetra acetic acid, an inhibitor of the previously described V , cholerae
	hemagglutinin/protease. When unnicked cholera enterotoxin was incubated with purified hemagglutinin/pro-
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tease, the unnicked A subunit was converted to a molecular weight consistent with that of the A₁ subunit as

10¹⁰/ml. Partially purified enterotoxin was analyzed by sodi-

um dodecvl sulfate-polyacrylamide gel electrophoresis with

12% polyacrylamide gels. A significant proportion of the A subunit was unnicked when flask-grown cultures with or

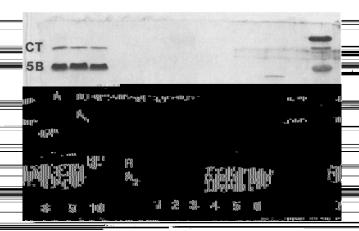
without EGTA (or 150 µg of Zincov per ml [MIC, 3 mg/ml]:

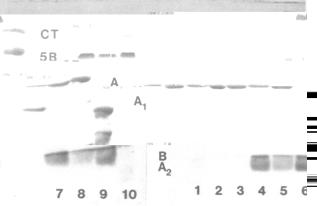
data not shown) were rapidly processed, i.e., within about 6

h (Fig. 1. lanes 8 and 9). When toxin was produced in

fermentor-grown cultures that were then processed at room

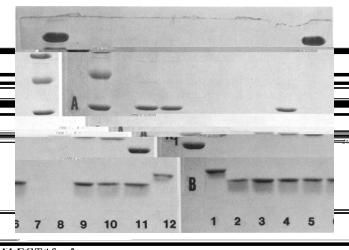
temperature during the next 2 days, the enterotoxin produced and processed in the absence of EGTA was almost





the culture grown in the presence of 6.6 mM EGIA and processed in the presence of 1 mM EGTA was almost completely unnicked (Fig. 2). Thus, the chelating agent protected the cholera toxin from being nicked. However,

when this unnicked, partially purified cholera toxin was left



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-	entration was estimated by a crude immunoassay, which is	peptide bonds upstream (12) from the cleavage site is an Arg-
		popular contact about carriers and the state
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<u>h</u>	iased by the simultaneous presence of choleragenoid (3.7).	Ser bond. a likely site of trypsin hydrolysis that would yield
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<u> </u>	· · · · · · · · · · · · · · · · · · ·	a slightly smaller A, fragment. It is also possible that other
	DISCUSSION	a slightly smaller A ₁ fragment. It is also possible that other proteases may effect activation by nicking neighboring pep-
	Cholera toxin is usually isolated in the nicked (activated)	tide bonds.
r. h	orm. Addition of EGTA. an inhibitor of HA/protease. can	In summary, although gut enzymes may play a contribu-
U	orm. Addition of EGTA, an inflibitor of HAzprotease, call	In Junimary, arthough fur only mey may a contribu-