U.S. PRODUCERS' QUESTIONNAIRE

GLYCINE FROM CHINA, INDIA, JAPAN, AND THAILAND

This questionnaire must be received by the Commission by <u>January 7, 2019</u>

See last page for filing instructions.

The information called for in this questionnaire is for use by the United States International Trade Commission in connection with countervailing duty investigations concerning glycine and certain precursor products from China, India, and Thailand and in connection with antidumping investigations concerning glycine and certain precursor products from India, Japan, and Thailand (Inv. Nos. 701-TA-603-605 and 731-TA-1413-1415 (Final)). The information requested in the questionnaire is requested under the authority of the Tariff Act of 1930, title VII. This report is mandatory and failure to reply as directed can result in a subpoena or other order to compel the submission of records or information in your firm's possession (19 U.S.C. § 1333(a)).

Name of firm

Δddrass

	State Zip Code
Website	
•	roduced or exported glycine and/or certain precursor products of glycine (as defined on next page) at anuary 1, 2015?
□ NO	(Sign the certification below and promptly return only this page of the questionnaire to the Commission)
☐ YES	(Complete all parts of the questionnaire, and return the entire questionnaire to the Commission)
•	ionnaire via the U.S. International Trade Commission <i>Drop Box</i> by clicking on the :: https://dropbox.usitc.gov/oinv/ . (PIN: GLYC)
	CERTIFICATION
ge and belief and of this certification	ation herein supplied in response to this questionnaire is complete and correct to the best of d understand that the information submitted is subject to audit and verification by the Commission on I also grant consent for the Commission, and its employees and contract personnel, to use
ge and belief and of this certification tion provided in to mission on the san odersigned, acknow of or other proce el (a) for develop and evaluations x 3; or (ii) by U.S.	d understand that the information submitted is subject to audit and verification by the Commission
ge and belief and if this certification ion provided in to mission on the san dersigned, acknow ing or other proce el (a) for develop and evaluations x 3; or (ii) by U.S.	d understand that the information submitted is subject to audit and verification by the Commission on I also grant consent for the Commission, and its employees and contract personnel, to use this questionnaire and throughout this proceeding in any other import-injury proceedings conducted time or similar merchandise. Towledge that information submitted in response to this request for information and throughout eedings may be disclosed to and used: (i) by the Commission, its employees and Offices, and containing or maintaining the records of this or a related proceeding, or (b) in internal investigations, and is relating to the programs, personnel, and operations of the Commission including under 5 U. In government employees and contract personnel, solely for cybersecurity purposes. I understand that an appropriate nondisclosure agreements
te and belief and for this certification on provided in the same of the same o	d understand that the information submitted is subject to audit and verification by the Commission on I also grant consent for the Commission, and its employees and contract personnel, to use this questionnaire and throughout this proceeding in any other import-injury proceedings conducted time or similar merchandise. Towledge that information submitted in response to this request for information and throughout eedings may be disclosed to and used: (i) by the Commission, its employees and Offices, and containing or maintaining the records of this or a related proceeding, or (b) in internal investigations, and is relating to the programs, personnel, and operations of the Commission including under 5 U. In government employees and contract personnel, solely for cybersecurity purposes. I understand that an appropriate nondisclosure agreements

PART I.—GENERAL INFORMATION

Background.--This proceeding was instituted in response to a petition filed on March 28, 2018, by GEO Specialty Chemical, Lafayette, Indiana and Chattem Chemicals, Inc., Chattanooga, Tennessee. Countervailing and antidumping duties may be assessed on the subject imports as a result of these proceedings if the Commission makes an affirmative determination of injury, threat, or material retardation, and if the U.S. Department of Commerce ("Commerce") makes an affirmative determination of subsidization and dumping. Questionnaires and other information pertinent to this proceeding are available

https://www.usitc.gov/investigations/701731/2018/glycine china india japan and thailand/final.htm.

<u>Glycine</u> covered by these investigations is glycine at any purity level or grade. This includes glycine of all purity levels, which covers all forms of crude or technical glycine including but not limited to sodium glycinate, glycine slurry and any other forms of amino acetic acid or glycine. Subject merchandise also includes glycine and precursors of dried crystalline glycine that are processed in a third country, including, but not limited to, refining or any other processing that would not otherwise remove the merchandise from the scope of the investigations if performed in the country of manufacture of the inscope glycine or precursors of dried crystalline glycine.

Glycine has the Chemical Abstracts Service (CAS) registry number of 56-40-6. Glycine and glycine slurry are classified under Harmonized Tariff Schedule of the United States (HTSUS) subheading 2922.49.43.00. Sodium glycinate is classified in the HTSUS under 2922.49.80.00. While the HTSUS subheadings and CAS registry number are provided for convenience and customs purposes, the written description of the scope of these investigations is dispositive.

Reporting of information.--If information is not readily available from your records, provide carefully prepared estimates. If your firm is completing more than one questionnaire (i.e., a producer, importer, and/or purchaser questionnaire), you need not respond to duplicated questions.

<u>Confidentiality</u>.--The commercial and financial data furnished in response to this questionnaire that reveal the individual operations of your firm will be treated as confidential by the Commission to the extent that such data are not otherwise available to the public and will not be disclosed except as may be required by law (see 19 U.S.C. § 1677f). Such confidential information will not be published in a manner that will reveal the individual operations of your firm; however, general characterizations of numerical business proprietary information (such as discussion of trends) will be treated as confidential business information only at the request of the submitter for good cause shown.

<u>Verification</u>.--The information submitted in this questionnaire is subject to audit and verification by the Commission. To facilitate possible verification of data, please keep all files, worksheets, and supporting documents used in the preparation of the questionnaire response. Please also retain a copy of the final document that you submit.

Release of information.--The information provided by your firm in response to this questionnaire, as well as any other business proprietary information submitted by your firm to the Commission in connection with this proceeding, may become subject to, and released under, the administrative protective order provisions of the Tariff Act of 1930 (19 U.S.C. § 1677f) and section 207.7 of the Commission's Rules of Practice and Procedure (19 CFR § 207.7). This means that certain lawyers and other authorized individuals may temporarily be given access to the information for use in connection with this proceeding or other import-injury proceedings conducted by the Commission on the same or similar merchandise; those individuals would be subject to severe penalties if the information were divulged to unauthorized individuals.

<u>D-GRIDS tool.</u>--The Commission has a tool that firms can use to move data from their own MS Excel compilation files into self-contained data tables within this MS Word questionnaire, thereby reducing the amount of cell-by-cell data entry that would be required to complete this form. This tool is a macroenabled MS Excel file available for download from the Commission's generic questionnaires webpage (https://www.usitc.gov/trade_remedy/question.htm) called the "D-GRIDs tool." Use of this tool to help your firm complete this questionnaire is *optional*. Firms opting to use the D-GRIDs tool to populate their data into this questionnaire will need the D-GRIDs specification sheet PDF file specific to this proceeding (available on the case page which is linked under the "Background" above) which includes the necessary references relating to this questionnaire, as well as the macro-enable MS Excel D-GRIDs tool itself from the generic questionnaires page. More detailed instructions on how to use the D-GRIDs tool are available within the D-GRIDs tool itself.

I-1a. <u>OMB statistics</u>.--Please report below the actual number of hours required and the cost to your firm of completing this questionnaire.

Hours	Dollars

The questions in this questionnaire have been reviewed with market participants to ensure that issues of concern are adequately addressed and that data requests are sufficient, meaningful, and as limited as possible. Public reporting burden for this questionnaire is estimated to average 50 hours per response, including the time for reviewing instructions, gathering data, and completing and reviewing the questionnaire.

We welcome comments regarding the accuracy of this burden estimate, suggestions for reducing the burden, and any suggestions for improving this questionnaire. Please attach such comments to your response or send to the Office of Investigations, USITC, 500 E St. SW, Washington, DC 20436.

I-1b.	TAA information release In the event that the U.S. International Trade Commission (USITC)
	makes an affirmative final determination in this proceeding, do you consent to the USITC's
	release of your contact information (company name, address, contact person, telephone
	number, email address) appearing on the front page of this questionnaire to the Departments of
	Commerce, Labor, and Agriculture, as applicable, so that your firm and its workers can be made
	eligible for benefits under the Trade Adjustment Assistance program?

Yes	No
1 5	1110

I-2. <u>Establishments covered</u>.--Provide the city, state, zip code, and brief description of each establishment covered by this questionnaire. If your firm is publicly traded, please specify the stock exchange and trading symbol in the footnote to the table. Firms operating more than one establishment should combine the data for all establishments into a single report.

"<u>Establishment</u>"--Each facility of a firm involved in the <u>production</u> of glycine, including auxiliary facilities operated in conjunction with (whether or not physically separate from) such facilities.

Establishments covered ¹	City, State	Zip (5 digit)	Description
1			
2			
3			
4			
5			
6			
¹ Additional discussion on establishments consolidated in this questionnaire:			

I-3. Petition support.--Does your firm support or oppose the petition?

Country	Support	Oppose	Take no position
China CVD			
India AD			
India CVD			
Japan AD			
Thailand AD			
Thailand CVD			

U.S. P	roducers' Questionnaire	Glycine (Final)	Page !	
I-4.	OwnershipIs your firm owned, in whole or in part, by any other firm?			
	□ No □ Ye	sList the following information.		
	Firm name	Address	Extent of ownership (percent)	
I-5.	foreign, that are engage United States or that a the United States?	ged in importing glycine from Chi	y related firms, either domestic or na, India, Japan and Thailand into the from China, India, Japan and Thailand to	
	Firm name	Country	Affiliation	

I-7.	•	elated producers Does your firm have any related firms, either domestic or foreign, that are ngaged in the production of glycine?			
	No YesList the	following information.			
	Firm name	Country	Affiliation		

PART II.--TRADE AND RELATED INFORMATION

Telephone

Further information on this part of the questionnaire can be obtained from Celia Feldpausch (202-205-2387, celia.feldpausch@usitc.gov). Supply all data requested on a celia.feldpausch@usitc.gov).

II-1.		nationPlease identify the responsible aff may contact that individual regardin	e individual and the manner by which ang the confidential information submitted
	Name		
	Title		
	Email		

II-2. <u>Changes in operations.</u>—Please indicate whether your firm has experienced any of the following changes in relation to the production of glycine since January 1, 2015.

(chec	k as many as appropriate)	(If checked, please describe; leave blank if not applicable)
	plant openings	
	plant closings	
	relocations	
	expansions	
	acquisitions	
	consolidations	
	prolonged shutdowns or production curtailments	
	revised labor agreements	
	other (e.g., technology)	

II-3a. **Production using same machinery.--**Please report your firm's production of products using the same equipment, machinery, or employees as used to produce glycine, and the combined production capacity on this shared equipment, machinery, or employees in the periods indicated.

"Nameplate capacity"— The maximum level of production that your establishment(s) could have obtained during the specified periods assuming maximum operating parameters and conditions (e.g., using equipment and machinery in place and ready to operate; twenty-four hours operating levels (hours per week/weeks per year) and no time for downtime, no maintenance, no repair, and no cleanup).

"Overall production capacity"— The level of production that your establishment(s) could reasonably have expected to attain during the specified periods. Assume normal operating conditions (i.e., using equipment and machinery in place and ready to operate; normal operating levels (hours per week/weeks per year) and time for downtime, maintenance, repair, and cleanup).

Note.--If your firm does not produce any out-of-scope merchandise on the same machinery and equipment as scope merchandise then the "overall production capacity" numbers reported in this question should be exactly equal to the "average production capacity" numbers reported in question II-7. If, however, your firm does produce out-of-scope merchandise using the same machinery and equipment as scope mercandhise, then the "average production capacity" reported in question II-7 should exclude the portion of "overall production capacity" that was used to produce this out-of-scope merchandise.

"Production" – All production in your U.S. establishment(s), including production consumed internally within your firm and production for another firm under a toll agreement.

Quantity (in 1,000 pounds)					
	Calendar years January –September				
Item	2015	2016	2017	2017	2018
Nameplate capacity					
Overall production capacity					
Production of: Glycine ¹	0	0	0	0	0
Other products ²					
Total	0	0	0	0	0

¹ Data entered for production of glycine will populate here once reported in question II-7.

II-3b. <u>Operating parameters.</u>—The overall production capacity reported in II-3a is based on the following operating parameters:

Hours per week	Weeks per year

² Please identify these products: _____

capacity	reported in	II-3a, and explain any changes in reported capacity.
	on constrai on capacity	ntsPlease describe the constraint(s) that set the limit(s) on your firm's
	shifting.—	
	-	able to switch production (capacity) between glycine and other products using uipment and/or labor?
No	Yes	If yes—(i.e., have produced other products or are able to produce other products) Please identify other actual or potential products:
b	etween pro	ribe the factors that affect your firm's ability to shift production capacity oducts (e.g., time, cost, relative price change, etc.), and the degree to which s enhance or constrain such shifts.
	Since Janua on of glycin	ary 1, 2015, has your firm been involved in a toll agreement regarding the e?
materials	and the se	agreement between two firms whereby the first firm furnishes the raw cond firm uses the raw materials to produce a product that it then returns a charge for processing costs, overhead, etc.
No	Yes	If yes Please describe the toll arrangement(s) and name the firm(s) involved.
	1 -	

U.S. Producers	' Questionnaire – Gl	ycine (Final)	
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II-5.	Foreign	trade	zones

(a) <u>Firm's FTZ operations</u>.--Does your firm produce glycine in and/or admit glycine into a foreign trade zone (FTZ)?

"Foreign trade zone" is a designated location in the United States where firms utilize special procedures that allow delayed or reduced customs duty payments on foreign merchandise. A foreign trade zone must be designed as such pursuant to the rules and procedures set forth in the Foreign-Trade Zones Act.

No	Yes	If yes Describe the nature of your firms operations in FTZs and identify the specific FTZ site(s).

(b) Other firms' FTZ operations.--To your knowledge, do any firms in the United States import glycine into a foreign trade zone (FTZ) for use in distribution of glycine and/or the production of downstream articles?

No	Yes	If yesIdentify the firms and the FTZs.

II-6. <u>Importer</u>.--Since January 1, 2015, has your firm imported glycine?

"Importer" – The person or firm primarily liable for the payment of any duties on the merchandise, or an authorized agent acting on his behalf. The importer may be the consignee, or the importer of record.

No	Yes	
		If yes <u>COMPLETE AND RETURN A U.S. IMPORTERS' QUESTIONNAIRE</u>

- II-7. **Production, shipment, and inventory data**.--Report your firm's production capacity, production, shipments, and inventories related to the production of glycine in its U.S. establishment(s) during the specified periods.
 - "Average production capacity" or "capacity" The level of production that your establishment(s) could reasonably have expected to attain during the specified periods. Assume normal operating conditions (i.e., using equipment and machinery in place and ready to operate; normal operating levels (hours per week/weeks per year) and time for downtime, maintenance, repair, and cleanup; and a typical or representative product mix).
 - "**Production**" All production in your U.S. establishment(s), including production consumed internally within your firm and production for another firm under a toll agreement.
 - "Commercial U.S. shipments" Shipments made within the United States as a result of an arm's length commercial transaction in the ordinary course of business. Report <u>net values</u> (i.e., gross sales values less all discounts, allowances, rebates, prepaid freight, and the value of returned goods) in U.S. dollars, f.o.b. your point of shipment.
 - "Internal consumption" Product consumed internally by your firm. Such transactions are valued at fair market value.
 - "Transfers to related firms" Shipments made to related domestic firms. Such transactions are valued at fair market value.
 - "Related firm" —A firm that your firm solely or jointly owns, manages, or otherwise controls.
 - "Export shipments" Shipments to destinations outside the United States, including shipments to related firms.
 - "Inventories" Finished goods inventory, not raw materials or work-in-progress.

Note: As requested in Part I of this questionnaire, please keep all supporting documents/records used in the preparation of the trade data, as Commission staff may contact your firm regarding questions on the trade data. The Commission may also request that your company submit copies of the supporting documents/records (such as production and sales schedules, inventory records, etc.) used to compile these data.

II-7. **Production, shipment, and inventory data.--Continued**

Quantity (in 1,000 pounds) and value (in \$1,000) Calendar years January –September							
				-			
Item	2015	2016	2017	2017	2018		
Average production capacity¹ (quantity) (A)							
Beginning-of-period inventories (quantity) (B)							
Production (quantity) Using HCN process (C)							
Using MCA process (D)							
Total production (E)	0	0		0 0	(
U.S. shipments: Commercial shipments: Quantity (F)							
Value (G)							
Internal consumption: ² Quantity (H)							
Value² (I)							
Transfers to related firms: ² Quantity (J)							
Value² (K)							
Export shipments: ³ Quantity (L)							
Value (M)							
End-of-period inventories (quantity) (N)							
¹ The production capacity reported is describe the methodology used to calcucapacity ² Internal consumption and transfers different basis for valuing these transac provide the data above at fair market value.	to related firm	n capacity, and	d explain any cl	hanges in reporto	ed vour firm use a		

<u>RECONCILIATION OF SHIPMENTS, PRODUCTION, AND INVENTORY.</u>--Generally, the data reported for the end-of-period inventories (i.e., line N) should be equal to the beginning-of-period inventories (i.e., line B), plus production (i.e., lines C and D), less total shipments (i.e., lines F, H, J and L). Please ensure that any differences are not due to data entry errors in completing this form, but rather reflect your firm's actual records; and, also provide explanations for any differences (e.g., theft, loss, damage, record systems issues, etc.) if they exist.

	Calendar years			January -September		
Reconciliation	2015	2016	2017	2017	2018	
B+C+D-F-H-J-L-N=						
should equal zero ("0") or						
provide an explanation.1	0	0	0	0	0	
¹ Explanation if the calculated fields al	¹ Explanation if the calculated fields above are returning values other than zero (i.e., "0") but are nonetheless					

accurate:_____.

II-8. <u>Channels of distribution</u>.--Report your firm's U.S. shipments (i.e., inclusive of commercial U.S. shipments, internal consumption, and transfers to related firms) by channel of distribution.

Quantity (in 1,000 pounds)							
	Calendar years			January –September			
Item	2015	2016	2017	2017	2018		
Channels of distribution: U.S. shipments: To distributors (O)							
To end users (P)							

<u>RECONCILIATION OF CHANNELS</u>.--Please ensure that the quantities reported for channels of distribution (i.e., lines O and P) in each time period equal the quantity reported for U.S. shipments (i.e., lines F, H, and J) in each time period. If the calculated fields below return values other than zero (i.e., "0"), the data reported must be revised prior to submission to the Commission.

	Calendar years			January –September		
Reconciliation	2015	2016	2017	2017	2018	
O + P - F - H - J = zero ("0"), if						
not revise.	0	0	0	0	0	

II-9. <u>U.S. shipments by product type</u>.--Report your firm's U.S. shipments (i.e., inclusive of commercial U.S. shipments, internal consumption, and transfers to related firms) by product type during the specified periods.

Quantity (in 1,000 pounds)						
	Calendar years			January –September		
Item	2015	2016	2017	2017	2018	
U.S. shipments: Technical grade glycine: Quantity (Q)						
Value (R)						
USP grade glycine: Quantity (S)						
Value (T)						
Pharmaceutical grade glycine, not injectable: Quantity (U)						
Value (V)						
Pharmaceutical grade glycine, injectable: Quantity (W)						
Value (X)						
In-scope glycine precursor products: Quantity (Y)						
Value (Z)						

<u>RECONCILIATION OF SHIPMENTS</u>.--Please ensure that the quantities and values reported for U.S. shipments by product type (i.e., lines Q through Y) in each time period equal the quantities and values reported for U.S. shipments (i.e., lines F through K) in each time period. If the calculated fields below return values other than zero (i.e., "0"), the data reported must be revised prior to submission to the Commission.

	Calendar years			January -September		
Reconciliation	2015	2016	2017	2017	2018	
Quantity: Q + S + U + W + Y - F - H - J = zero ("0"), if not						
revise.	0	0	0	0	0	
Value: $R + T + V + X + Z - G - I - K$ = zero ("0"), if not revise.	0	0	0	0	0	

II-10. <u>U.S. shipments by certification</u>.--Report your firm's U.S. shipments (i.e., inclusive of commercial U.S. shipments, internal consumption, and transfers to related firms) by certification in 2017.

U.S. shipments	Calendar year 2017 Quantity (1,000 pounds)
FDA certified, but not EDQM certified (AA)	
EDQM certified, but not FDA certified (AB)	
Both FDA and EDQM certified (AC)	
Neither FDA nor EDQM certified (AD)	

<u>RECONCILIATION OF SHIPMENTS</u>.--Please ensure that the quantities reported for U.S. shipments by certification (i.e., lines AA through AD) in each time period equal the quantities reported for U.S. shipments in 2017 (i.e., lines D, F and H) reported in question II-7. If the calculated fields below return values other than zero (i.e., "0"), the data reported must be revised prior to submission to the Commission.

Reconciliation	Calendar year 2017
Quantity: $AA + AB + AC + AD - D - F - H = zero$	
("0"), if not revise.	0

Explanation of trends:

II-11. <u>Employment data</u>.--Report your firm's employment-related data related to the production of glycine and provide an explanation for any trends in these data.

"Production and Related Workers" (PRWs) includes working supervisors and all nonsupervisory workers (including group leaders and trainees) engaged in fabricating, processing, assembling, inspecting, receiving, storage, handling, packing, warehousing, shipping, trucking, hauling, maintenance, repair, janitorial and guard services, product development, auxiliary production for plant's own use (e.g., power plant), recordkeeping, and other services closely associated with the above production operations.

Average number employed may be computed by adding the number of employees, both full time and part time, for the 12 pay periods ending closest to the 15th of the month and divide that total by 12. For the January to September periods, calculate similarly and divide by 9.

"Hours worked" includes time paid for sick leave, holidays, and vacation time. Include overtime hours actually worked; do not convert overtime pay to its equivalent in straight time hours.

"Wages paid" – Total wages paid before deductions of any kind (e.g., withholding taxes, old-age and unemployment insurance, group insurance, union dues, bonds, etc.). Include wages paid directly by your firm for overtime, holidays, vacations, and sick leave.

		Calendar years	S	January –	September
Item	2015	2016	2017	2017	2018
Average number of PRWs (number)					
Hours worked by PRWs (1,000 hours)					
Wages paid to PRWs (\$1,000)					

	Related firms.—If your firm reported transfers to related firms in question II-7, please indicate the nature of the relationship between your firm and the related firms (e.g., joint venture, wholly owned subsidiary), whether the transfers were priced at market value or by a non-market formula, whether your firm retained marketing rights to all transfers, and whether the related firms also processed inputs from sources other than your firm.
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U.S. Producers'	Questionnaire – Glycine	(Final)

II-12. <u>Purchases</u>.--Other than imports, has your firm otherwise purchased glycine (domestic or imported) any time since January 1, 2015?

"Purchase" – A transaction to buy product from a U.S. corporate entity such as another U.S. producer, a U.S. distributor, or a U.S. firm that has directly imported the product.

"Import" –A transaction to buy from a foreign supplier where your firm is the importer of record.

No	If yes Report such purchases below and explain the reasons for your firms' purchases:

	(Quanti	ity <i>in 1,000 po</i>	unds)		
	(Calendar years	S	January -	-September
Item	2015	2016	2017	2017	2018
Purchases from U.S. importers ¹ of glycine from— China					
India					
Japan					
Thailand					
All other sources					
Purchases from domestic producers ²					
Purchases from other sources ²					

ittei	r by sou	rce, please identify the s	source for each listed supplier:
2	Please I	ist the name of the prod	ducer(s) or U.S. distributor(s) from which your firm purchased this product:
	11 12	Other evalenations	If your firm would like to further evaluing response to a question in Port II

II-13.	that did not provide a narrative box, please note the question number and the explanation in the space provided below. Please also use this space to highlight any issues your firm had in providing the data in this section, including but not limited to technical issues with the MS Word questionnaire.
	questionnaire.

PART III.--FINANCIAL INFORMATION

Address questions on this part of the questionnaire to Jennifer Brinckhaus (202-205-3188, jennifer.brinckhaus@usitc.gov).

Name	
Title	
Email	
Telephone	
Accounting sy	<u>ystem</u> Briefly describe your firm's financial accounting system.
A.	When does your firm's fiscal year end (month and day)?
	If your firm's fiscal year changed during the data-collection period, explain below:
	NotePlease note that we are requesting that firms report their financial day on a calendar year basis.
B.1.	Describe the lowest level of operations (e.g., plant, division, company-wide) which financial statements are prepared that include glycine:
2.	Does your firm prepare profit/loss statements for glycine:
3.	How often did your firm (or parent company) prepare financial statements
	(including annual reports, 10Ks)? Please check relevant items below. ☐ Audited, ☐ unaudited, ☐ annual reports, ☐ 10Ks, ☐ 10 Qs, ☐ Monthly, ☐ quarterly, ☐ semi-annually, ☐ annually
4.	Accounting basis: GAAP, cash, tax, or other comprehen

Note: As requested in Part I of this questionnaire, please keep all supporting documents/records used in the preparation of the financial data, as Commission staff may contact your firm regarding questions on the financial data. The Commission may also request that your company submit copies of the supporting documents/records (financial statements, including internal profit-and-loss statements for the division or product group that includes glycine, as well as specific statements and worksheets) used to compile these data.

U.S. Producers' Questionnaire - Glycine (Fina	U.S.	Producers'	Questionnaire -	Glycine	(Fina
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Allocatio	on basisBriefly describe your firm's allocation basis, if any, for COGS, SG&A, and
	expense and other income and expenses.

III-5. **Product listing**.--Please list the products your firm produced in the facilities in which your firm produced glycine, and provide the share of net sales accounted for by these products in 2017.

Products	Share of sales
Glycine	%
	%
	%
	%
	%

Yes	Continue to ques	ion III-7	NoContinue to	question III-9a.	
nputs fro	om related supplier	s detailedPl	ease identify the input	ts used in the production	
				are reflected in question	
	·	•	-	ant input on the basis of	
	most recently completed fiscal year. For "Input valuation" please describe the basis, as				
recorded in your company's own accounting system, of the purchase cost from the rela					
			•		
upplier;	e.g., the related sup	plier's actual	ng system, of the purch cost, cost plus, negotia		
upplier;		plier's actual	•		
supplier;	e.g., the related sup	plier's actual	cost, cost plus, negotia		
upplier; o	e.g., the related sup	plier's actual e.	cost, cost plus, negotia	ated transfer price to	
upplier; (pproxim	e.g., the related sup	plier's actual e.	cost, cost plus, negotia	ated transfer price to	
upplier; (pproxim	e.g., the related sup	plier's actual e.	cost, cost plus, negotia	ated transfer price to	
supplier; on the supproxim supproxim supproxim supproximal supproximals supproximately supproxim	e.g., the related sup ate fair market valu	Related su	ost, cost plus, negotia	Share of total CO	
supplier; of approxim	e.g., the related sup ate fair market valu	Related su	cost, cost plus, negotia	Share of total CO	
supplier; on the supproxim supproxim supproxim supproximal supproximals supproximately supproxim	e.g., the related sup ate fair market valu	Related su	ost, cost plus, negotia	Share of total CO	
upplier; opproxim Input Input va	e.g., the related sup ate fair market valu	Related su	ipplier accounting books and	Share of total CO	
Input Input va	e.g., the related sup ate fair market valu luation as recorded rchased from relat	Related su I in the firm's ed suppliers d in III-7, were	upplier accounting books and	Share of total Co	

III-9a. Operations on glycine.--Report the revenue and related cost information requested below on the glycine operations of your firm's U.S. establishment(s).¹ Do not report resales of products. Note that internal consumption and transfers to related firms must be valued at fair market value. Input purchases from related suppliers should be consistent with and based on information in the firm's accounting books and records. Provide data for your firm on a calendar year basis, and for the specified interim periods.

	С	alendar years	January-Sep	otember	
ltem	2015	2016	2017	2017	2018
Net sales quantities: ²					
Commercial sales ("CS")					
Internal consumption ("IC")					
Transfers to related firms ("Transfers")					
Total net sales quantities	0	0	0	0	C
Net sales values: ² Commercial sales					
Internal consumption					
Transfers to related firms					
Total net sales values	0	0	0	0	С
Cost of goods sold (COGS): ³ Raw materials					
Direct labor					
Other factory costs					
Total COGS	0	0	0	0	C
Gross profit or (loss)	0	0	0	0	С
Selling, general, and administrative (SG&A) expenses: Selling expenses					
General and administrative expenses					
Legal expenses ⁴					
Total SG&A expenses	0	0	0	0	0
Operating income (loss)	0	0	0	0	0
Other expenses and income: Interest expense					
All other expense items					
All other income items					
Net income or (loss) before income taxes	0	0	0	0	C
Depreciation/amortization included above					

¹ Include only sales (whether <u>domestic or export</u>) and costs related to your <u>U.S. manufacturing operations</u>.

Note -- The table above contains calculations that will appear when you have entered data in the MS Word form fields.

² Less discounts, returns, allowances, and prepaid freight. The quantities and values should approximate the corresponding shipment quantities and values reported in Part II of this questionnaire.

³ COGS (whether for domestic or export sales) should include costs associated with CS, IC, and Transfers.

⁴ Please indicate where these legal expenses appear in the company's own books and records (e.g., within G&A, other expenses, etc.):

III-9b.	<u>Financial data reconciliation</u> The calculable line items from question III-9a (i.e., total net sales
	quantities and values, total COGS, gross profit (or loss), total SG&A, and net income (or loss))
	have been calculated from the data submitted in the other line items. Do the calculated fields
	return the correct data according to your firm's financial records ignoring non-material
	differences that may arise due to rounding?

Yes	No	If no If the calculated fields do not show the correct data, please double check the feeder data for data entry errors and revise. Also, check signs accorded to the post operating income line items; the two expense line items should report positive numbers (i.e., expenses are positive and incomes or reversals are negativeinstances of the latter should be rare in those lines) while the income line item also in most instances should have its value be a positive number (i.e., income is positive, expenses or reversals are negative). If after reviewing and potentially revising the feeder data your firm has provided, the differences between your records and the calculated fields persist please identify and discuss the differences in the space below.

III-9c. Raw materials for producers using the Strecker amino acid synthesis (HCN) process.--If your firm produces glycine using the Strecker amino acid synthesis process, please report the raw material costs (reported in III-9a) for the following raw material inputs:

Value (<i>in \$1,000</i>)							
	Calendar years			January-September			
Input	2015	2016	2017	2017	2018		
HCN process raw materials: Formaldehyde							
Hydrogen cyanide							
Ammonia							
Other material inputs ¹							
¹ Please indicate any other notab	e "other" raw r	naterials not ex	pressly identific	ed above:			

III-9d. Raw materials for producers using the monochloroacetic acid (MCAA) process.--If your firm produces glycine using the monochloroacetic acid process, please report the raw material costs (reported in III-9a) for the following raw material inputs:

Value (in \$1,000)						
	Calendar years			January-September		
Input	2015	2016	2017	2017	2018	
MCAA process raw materials: Ammonia						
Monochloroacetic acid						
Hexamethylenetetramine or other catalyst(s)						
Other material inputs ¹						

III-10. Nonrecurring items (charges and gains) included in the subject product financial results.--For each annual and interim period for which financial results are reported in question III-9a, please specify all material (significant) nonrecurring items (charges and gains) in the schedule below, the specific question III-9a line item where the nonrecurring items are included, a brief description of the relevant nonrecurring items, and the associated values (in \$1,000), as reflected in question III-9a; i.e., if an aggregate nonrecurring item has been allocated to question III-9a, only the allocated value amount included in question III-9a should be reported in the schedule below. Note: The Commission's objective here is to gather information only on material (significant) nonrecurring items which impacted the reported financial results of the subject product in question III-9a.

		Calendar years		January-Septembe			
	2015	2016	2017	2017	2018		
Item	Value (<i>\$1,000</i>)						
Nonrecurring item 1							
Nonrecurring item 2							
Nonrecurring item 3							
Nonrecurring item 4							
Nonrecurring item 5							
Nonrecurring item 6							
Nonrecurring item 7							

Nonrecurring item: In this table please provide a brief description of each nonrecurring item reported above and indicate the specific line item in table III-9a where the nonrecurring item is classified.

	Description of the nonrecurring item	Income statement classification of the nonrecurring item
Nonrecurring item 1		
Nonrecurring item 2		
Nonrecurring item 3		
Nonrecurring item 4		
Nonrecurring item 5		
Nonrecurring item 6		
Nonrecurring item 7		

III-11.	Classification of identified nonrecurring items (charges and gains) in the accounting books and records of the company.—If non-recurring items were reported in question III-10 above, please identify where your company recorded these items in your accounting books and records in the normal course of business; i.e., just as responses to question III-10 identify where these items are reported in question III-9a.

III-12. <u>Asset values</u>.--Report the <u>total</u> assets (i.e., both current and long-term assets) associated with the production, warehousing, and sale of glycine. If your firm does not maintain some or all of the specific asset information necessary to calculate total assets for glycine in the normal course of business, please estimate this information based upon a method (such as production, sales, or costs) that is consistent with relevant cost allocations in question III-9a. Provide data on a calendar year basis.

Note: Total assets should reflect <u>net assets</u> after any accumulated depreciation and allowances deducted.

Total assets should be <u>allocated to the subject products</u> if these assets are also related to other products. Please provide a <u>brief explanation if there are any substantial changes</u> in total asset value during the period; e.g., due to asset write-offs, revaluation, and major purchases.

Value (<i>in \$1,000</i>)					
	Calendar years 2015 2016 2017				
Item					
Total assets (net) ¹					
¹ Describe					

III-13. <u>Capital expenditures and research and development expenses</u>.--Report your firm's capital expenditures and research and development expenses for glycine. Provide data for your firm on a calendar year basis, and for the specified interim periods.

Value (in \$1,000)					
	Calendar years January-September				
ltem	2015	2016	2017	2017	2018
Capital expenditures ¹					
Research and development expenses ²					

¹ Please describe the nature, focus, and significance of your firm's capital expenditures on the subject product.

² Please describe the nature, focus, and significance of your firm's R&D expenses related to subject product.

III-14. <u>Data consistency and reconciliation</u>.--Please confirm that your firm's financial data for questions III-9a, 12, and 13 are based on a calendar year basis. The quantities and values reported in question III-9a should reconcile with the data reported in question II-7 (including export shipments).

RECONCILIATION OF TRADE VS FINANCIAL DATA.--Please ensure that the quantities and values reported for total shipments in part II equal the quantities and values reported for total net sales in part III of this questionnaire in each time period unless the financial data from part III are reported on a fiscal year basis, in which case only the interim periods must reconcile. If the calculated fields below return values other than zero (i.e., "0") and both are being reported on a calendar basis, please explain the discrepancy below.

		Calendar years		January-S	eptember
Reconciliation	2015	2016	2017	2017	2018
Quantity: Trade data from question II-7 (lines F, H, J and L) less financial total net sales quantity data from question III-9a, = zero ("0").	0	0	0	0	0
Value: Trade data from question II-7 (lines G, I, K and M) less financial total net sales value data from question III-9a, = zero ("0").	0	0	0	0	0

Do these data in question III-9a reconcile with data in question II-7?

Yes	No	If no, please explain.

If your responses to any of the items in questions III-15, III-16, and III-17 differ by country, please describe these differences and, as applicable, indicate which country or countries your response refers to in the relevant form fields.

III-15. <u>Effects of imports on investment</u>.--Since January 1, 2015, has your firm experienced any actual negative effects on its return on investment or the scale of capital investments as a result of imports of glycine from China, India, Japan, or Thailand?

No	Yes		
		If yes, my firm has experier	nced actual negative effects as follows:
	(checi	k as many as appropriate)	(please describe)
		Cancellation, postponement, or rejection of expansion projects	
		Denial or rejection of investment proposal	
		Reduction in the size of capital investments	
		Return on specific investments negatively impacted	
		Other	

III-16.	Effects of imports on growth and developmentSince January 1, 2015, has your firm
	experienced any actual negative effects on its growth, ability to raise capital, or existing
	development and production efforts (including efforts to develop a derivative or more advanced
	version of the product) as a result of imports of glycine from China, India, Japan, or Thailand?

	-		
No	Yes		
		If yes, my firm has experie	nced actual negative effects as follows.
	(chec	ck as many as appropriate)	(please describe)
		Rejection of bank loans	
		Lowering of credit rating	
		Problem related to the issue of stocks or bonds	
		Ability to service debt	
		Other	

U.S. Producers'	Questionnaire –	Glycine (Final)

III-17.	Anticipated effects of importsDoes your firm anticipate any negative effects due to imports of
	glycine from China, India, Japan, or Thailand?

No	Yes	If yes, my firm anticipates negative effects as follows:

III-18.	Other explanations.—If your firm would like to further explain a response to a question in Part III that did not provide a narrative box, please note the question number and the explanation in the space provided below. Please also use this space to highlight any issues your firm had in providing the data in this section, including but not limited to technical issues with the MS Word questionnaire.

PART IV.--PRICING AND MARKET FACTORS

Further information on this part of the questionnaire can be obtained from Nabil Abbyad (Nabil.Abbyad@usitc.gov, 202-708-1446).

IV-1. <u>Contact information</u>.--Please identify the individual that Commission staff may contact regarding the confidential information submitted in part IV.

Name	
Title	
Email	
Telephone	

PRICE DATA

- IV-2. This question requests quarterly quantity and value data for your firm's commercial shipments to unrelated U.S. customers since January 1, 2015 of the following products produced by your firm.
 - <u>Product 1</u>.--Pharmaceutical-grade glycine -- a white, odorless, crystalline powder with a sweet taste, having an assay (glycine content) of 98.5 percent to 101.5 percent (dry basis), and \leq 7ppm chloride, \leq 65 ppm sulfate, and \leq 1 ppm heavy metals.
 - <u>Product 2</u>.--USP-grade glycine -- a white, odorless, crystalline powder with a sweet taste, having an assay (glycine content) of 98.5 percent to 101.5 percent (dry basis) and \leq 70 ppm chloride, \leq 65 ppm sulfate, \leq 20 ppm heavy metals, and not otherwise qualifying as pharmaceutical-grade glycine.
 - <u>Product 3.</u>--Technical-grade glycine -- a white, off-white, or slightly yellow crystalline powder, having an assay (glycine content) of 98.5 percent to 101.5 percent (dry basis), with maximum chlorides of 0.4 percent, and not otherwise qualifying as USP-grade glycine.

Please note that values should be <u>f.o.b.</u>, <u>U.S.</u> point of shipment and should not include U.S.-inland transportation costs. Values should reflect the *final net* amount paid to your firm (i.e., should be net of all deductions for discounts or rebates in the quarter of the original sale).

IV-2 (a). During January 2015-September 2018, did your firm produce and sell to unrelated U.S. customers any of the above listed products (or any products that were competitive with these products)?

YesPlease complete the following pricing data table as appropriate.
NoSkip to question IV-3.

IV-2(b). Price data.--Report below the quarterly price data¹ for pricing products² produced and sold by your firm.

Report data in <u>actual pounds</u> (not 1,000s) and <u>actual dollars</u> (not 1,000s).

(Quantity in pounds, value in dollars)						
	Product 1		Produ	ıct 2	Produ	ıct 3
Period of shipment	Quantity	Value	Quantity	Value	Quantity	Value
2015:						
January-March						
April-June						
July-September						
October-December						
2016:						
January-March						
April-June						
July-September						
October-December						
2017:						
January-March						
April-June						
July-September						
October-December						
2018:						
January-March						
April-June						
July-September						

Note. —If your firm's product does not exactly	y meet the product specifications but is competitive with the specified product,
provide a description of your firm's product.	Also, please explain any anomalies in your firm's reported pricing data.

	provide a description of your firm's product. Also, please explain any anomalies in your firm's reported pricing data.
	Product 1:
	Product 2:
	Product 3:
ı	

² Pricing product definitions are provided on the first page of Part IV.

IV-2 (c). Price data checklist	-Please check that the pricing data in question IV-	-2(b) has been co	rrectly
reported.			

Is the price data reported above:	√ if Yes
In actual dollars (not \$1,000) and actual pounds (not 1,000 pounds)?	
F.o.b. U.S. point of shipment (i.e., does not include U.S. transport costs)?	
Net of all discounts and rebates?	
Have returns credited to the quarter in which the sale occurred?	
Less than reported commercial shipments in question II-7 in each year?	

	Less than reported commercial simplification in 7 in each year.		
IV-2 (d).	Pricing data methodologyPlease describe the method and the kinds of docume	nts/records	
	hat were used to compile your price data.	,	
Г			

Note: As requested in Part I of this questionnaire, please keep all supporting documents/records used in the preparation of the price data, as Commission staff may contact your firm regarding questions on the price data. The Commission may also request that your company submit copies of the supporting documents/records (such as sales journal, invoices, etc.) used to compile these data.

				-
110	Duadinages	Questionnaire -		/F:l
11 >	Producers	THESTIONNAIRE -	GIVCIDE	IFINAL

IV-3.	<u>Price setting.</u> How does your firm determine the prices that it charges for sales of glycine
	(check all that apply)? If your firm issues price lists, please submit sample pages of a recent list.

Transaction by transaction	Contracts	Set price lists	Other	If other, describe

IV-4. <u>Discount policy</u>.--Please indicate and describe your firm's discount policies (*check all that apply*).

Quantity discounts	Annual total volume discounts	No discount policy	Other	Describe

IV-5. Pricing terms.--

(a) What are your firm's typical sales terms for its U.S.-produced glycine?

Net 30 days	Net 60 days	2/10 net 30 days	Other	Other (specify)

(b) On what basis are your firm's prices of domestic glycine usually quoted (check one)?

Delivered	F.o.b.	If f.o.b., specify point

IV-6. <u>Contract versus spot.</u>--Approximately what share of your firm's sales of its U.S.-produced glycine in 2017 was on a (1) long-term contract basis, (2) annual contract basis, (3) short-term contract basis, and (4) spot sales basis?

	Type of sale					
ltem	Short-term contracts (multiple deliveries for less than 12 months)	Annual contracts (multiple deliveries for 12 months)	Long-term contracts (multiple deliveries for more than 12 months)	Spot sales (for a single delivery)	Total (shoul sum to 100.0%	d o
Share of 2017 sales	%	%	%	%	0.0	%

IV-7. <u>Contract provisions.</u>--Please fill out the table regarding your firm's typical sales contracts for U.S.-produced glycine (or check "not applicable" if your firm does not sell on a long-term, short-term and/or annual contract basis).

Typical sales contract provisions	ltem	Short-term contracts (multiple deliveries for less than 12 months)	Annual contracts (multiple deliveries for 12 months)	Long-term contracts (multiple deliveries for more than 12 months)
Average contract duration	No. of days		365	
Price renegotiation	Yes			
(during contract period)	No			
	Quantity			
Fixed quantity and/or price	Price			
aa, e. pee	Both			
Meet or release	Yes			
provision	No			
Not applicab	le			

IV-8. <u>Lead times.</u>—What is your firm's share of sales from inventory and produced to order and what is the typical lead time between a customer's order and the date of delivery for your firm's sales of its U.S.-produced glycine?

Source	Share of 2017 sales	Lead time (Average number of days)
From inventory	%	
Produced to order	%	
Total (should sum to 100.0%)	0.0 %	

	IV-9.	Shipping	information
--	-------	----------	-------------

(a)	What is the approximate percentage of the cost of U.Sproduced glycine that i
	accounted for by U.S. inland transportation costs? percent
71.3	NAMES and apply a supply and the transport of the state o

(b) Who generally arranges the transportation to your firm's customers' locations?

Your firm Purchaser (check one)

(c) Indicate the approximate percentage of your firm's sales of glycine that are delivered the following distances from its production facility.

Distance from production facility	Share
Within 100 miles	%
101 to 1,000 miles	%
Over 1,000 miles	%
Total (should sum to 100.0%)	0.0 %

IV-10. **Geographical shipments.--**In which U.S. geographic market area(s) has your firm sold its U.S.-produced glycine since January 1, 2015 (check all that apply)?

Geographic area	√ if applicable
NortheastCT, ME, MA, NH, NJ, NY, PA, RI, and VT.	
Midwest.–IL, IN, IA, KS, MI, MN, MO, NE, ND, OH, SD, and WI.	
Southeast.—AL, DE, DC, FL, GA, KY, MD, MS, NC, SC, TN, VA, and WV.	
Central Southwest.—AR, LA, OK, and TX.	
Mountains.–AZ, CO, ID, MT, NV, NM, UT, and WY.	
Pacific Coast.—CA, OR, and WA.	
Other .—All other markets in the United States not previously listed, including AK, HI, PR, and VI.	

IV-11. <u>End uses.--</u>List the end uses of the glycine that your firm manufactures. For each end-use product, what percentage of the <u>total cost</u> is accounted for by glycine and other inputs?

		t of end use product ted for by	Total
End use product	Glycine	Other inputs	(should sum to 100.0% across)
	%	%	0.0 %
	%	%	0.0 %
	%	%	0.0 %

IV-12.	Pharmaceutical-grade glycine	
	i marmaccatical grade gryeme	•

(a)	Did your firm produ	ce pharmaceutical-grade glycine since 2015?
	☐ No	YesPlease complete the table below.
(b)	Please describe the accounted for by th	packaging type(s) and estimate the percentage of the $\underline{\text{total cost}}$ is e packaging.

Packaging type	Packaging (percent of total cost)
	%
	%
	%

(c) Please describe the pharmaceutical end use(s) and estimate the average price premium of these products when compared to USP-grade glycine prices.

Pharmaceutical end use	Price premium compared to USP-grade (percent)
	%
	%
	%

		No		YesPlea	se fill out t	he tak	ole.	
			End u	ıse in wh	ich this	На		anges in the price of this substitu ffected the price for glycine?
	Substitute	2	sub	stitute is	used	No	Yes	Explanation
1.								
2.								
3. IV-1	States (if I	known) f	or glycine	has char		lanuar nanges	y 1, 2 s in de	States and outside of the United 015. Explain any trends and descremand.
	States (if I the princi	known) f	or glycine	has char	ged since	lanuar nanges Fluc	γ1,2	015. Explain any trends and descr
	States (if I	known) f	or glycine rs that ha	has char ve affecto No	nged since and these characteristics of these characteristics.	lanuar nanges Fluct wit	y 1, 2 s in de tuate h no	015. Explain any trends and descremand.
IV-1	States (if I the princi	known) f pal facto	or glycine rs that ha Overall	has char ve affecto No	nged since and these ch	lanuar nanges Fluct wit	y 1, 2 s in de tuate h no	015. Explain any trends and descremand.
IV-1	States (if I the princi	known) f pal facto	or glycine rs that ha Overall	has char ve affecto No	nged since and these ch	lanuar nanges Fluct wit	y 1, 2 s in de tuate h no	015. Explain any trends and descremand.
IV-1	States (if I the princi Market the United State the United State	known) f pal facto ates ates hanges	Overall increase	No change	Overall decrease	Fluct wit clear	tuate h no trenc	015. Explain any trends and descremand.
IV-1	States (if I the princi	known) f pal facto ates ates hanges	Overall increase -Have the ycine since	No change	Overall decrease	Fluct wit clear	tuate h no trenc	O15. Explain any trends and descremand. Explanation and factors in the product range, product mi

IV-16.	Conditions	of com	petition

(a)	Is the glycine market subject to business cycles (other than general economy-wide
	conditions) and/or other conditions of competition distinctive to glycine? If yes, describe.

	Check all t	hat apply	/ .	F	Please describe.
	No Yes-Business cycles (e.g. seasonal business) Yes-Other distinctive conditions of competition			S	Skip to question IV-17.
	(b) If yes, have there been any changes glycine since January 1, 2015?				n the business cycles or conditions of competition for
	No	Yes	If yes, o	lescribe.	
	January 1, declining to	2015 (exa o accept r	mples inclunew custom	ude placing coners or renew	declined, or been unable to supply glycine since ustomers on allocation or "controlled order entry," v existing customers, delivering less than the mely shipment commitments, etc.)?
	No	Yes	If yes, plea	se describe.	
IV-18.	Raw mater	r <mark>ials</mark> Hov	w have glyc	ine raw mate	erial prices changed since January 1, 2015?
	Overall	No	Overall	Fluctuate with no	Explain, noting how raw material price changes

IV-19. **Product quality issues.--**Have any of your customers returned glycine or canceled orders due to quality issues such as product impurities or caking issues, since January 1, 2015?

No	Yes	If yes, please describe.

IV-20. <u>Interchangeability</u>.--Is glycine produced in the United States and in other countries interchangeable (i.e., can they physically be used in the same applications)?

Please indicate A, F, S, N, or 0 in the table below:

- A = the products from a specified country-pair are *always* interchangeable
- F = the products are *frequently* interchangeable
- S = the products are *sometimes* interchangeable
- N = the products are *never* interchangeable
- 0 = *no familiarity* with products from a specified country-pair

Country-pair	China	India	Japan	Thailand	Other countries
United States					
China					
India					
Japan					
Thailand					
For any country-pair producing glycine that is <i>sometimes</i> or <i>never</i> interchangeable, identify the country-pair and explain the factors that limit or preclude interchangeable use:					

IV-21. <u>Factors other than price</u>.--Are differences other than price (e.g., quality, availability, transportation network, product range, technical support, *etc*.) between glycine produced in the United States and in other countries a significant factor in your firm's sales of the products?

Please indicate A, F, S, N, or 0 in the table below:

A = such differences are *always* significant

F = such differences are *frequently* significant

S = such differences are *sometimes* significant

N = such differences are *never* significant

0 = no familiarity with products from a specified country-pair

Country-pair	China	India	Japan	Thailand	Other countries
United States					
China					
India					
Japan					
Thailand					

For any country-pair for which factors other than price *always* or *frequently* are a significant factor in your firm's sales of glycine, identify the country-pair and report the advantages or disadvantages imparted by such factors:

IV-22. <u>Customer identification</u>.--List the names and contact information for your firm's 10 largest U.S. customers for glycine since January 1, 2015. Indicate the share of the quantity of your firm's total shipments of glycine that each of these customers accounted for in 2017.

	Customer's name	City	State	Share of 2017 sales (%)
1				
2				
3				
4				
5				
6				
7				
8				
9				
10				

IV-23. Competition from imports

(a) <u>Lost revenue</u>.--Since January 1, 2015: To avoid losing sales to competitors selling glycine from China, India, Japan, and/or Thailand, did your firm:

Item	No	Yes
Reduce prices		
Roll back announced price increases		

(b) <u>Lost sales.</u>--Since January 1, 2015: Did your firm lose sales of glycine to imports of this product from China, India, Japan, and/or Thailand?

No	Yes

IV-24.	Other explanationsIf your firm would like to further explain a response to a question in Part
	IV that did not provide a narrative response box, please note the question number and the
	explanation in the space provided below. Please also use this space to highlight any issues your
	firm had in providing the data in this section, including but not limited to technical issues with
	the MS Word questionnaire.

HOW TO FILE YOUR QUESTIONNAIRE RESPONSE

This questionnaire is available as a "fillable" form in MS Word format on the Commission's website at: LINK

Please do not attempt to modify the format or permissions of the questionnaire document. Please submit the completed questionnaire using one of the methods noted below. If your firm is unable to complete the MS Word questionnaire or cannot use one of the electronic methods of submission, please contact the Commission for further instructions.

• <u>Upload via Secure Drop Box</u>.—Upload the MS Word questionnaire along with a scanned copy of the signed certification page (page 1) through the Commission's secure upload facility:

Web address: https://dropbox.usitc.gov/oinv/ Pin: GLYC

• E-mail.—E-mail the MS Word questionnaire to celia.feldpausch@usitc.gov; include a scanned copy of the signed certification page (page 1). Submitters are strongly encouraged to encrypt nonpublic documents that are electronically transmitted to the Commission to protect your sensitive information from unauthorized disclosure. The USITC secure drop-box system and the Electronic Document Information System (EDIS) use Federal Information Processing Standards (FIPS) 140-2 cryptographic algorithms to encrypt data in transit. Submitting your nonpublic documents by a means that does not use these encryption algorithms (such as by email) may subject your firm's nonpublic information to unauthorized disclosure during transmission. If you choose a non-encrypted method of electronic transmission, the Commission warns you that the risk of such possible unauthorized disclosure is assumed by you and not by the Commission.

If your firm does not produce this product, please fill out page 1, print, sign, and submit a scanned copy to the Commission.

<u>Parties to this proceeding</u>.—If your firm is a party to this proceeding, it is required to serve a copy of the completed questionnaire on parties to the proceeding that are subject to administrative protective order (see 19 CFR § 207.7). A list of such parties may be obtained from the Commission's Secretary (202-205-1803). A certificate of service must accompany the completed questionnaire you submit (see 19 CFR § 207.7). Service of the questionnaire must be made in paper form.