DEPARTMENT OF HEALTH AND HUMAN SERVICES FOOD AND DRUG ADMINISTRATION			
DISTRICT OFFICE ADDRESS AND PHONE NUMBER		DATE(S) OF INSPECTION	
12420 Parklawn Drive, Room 2032 Rockville, MD 20857		10/22-26/2018	
		FEI NUMBER	
Industry Information: www.fda.gov/oc/industry		3004161218	
NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT IS ISSUED			
TO: Janmejay R. Vyas, Chairman			
RRM NAME STREET ADDRESS			
Dishman Carbogen Amcis Ltd	Survey No. 47, Paiki Sub Plot No. 1, Lodariyal Sanand-Bavla		
CITY, STATE AND ZIP CODE	TYPE OF ESTABLISHMENT INSPECTED		
Ahmedabad 382 220, Gujarat, India	API Manufacturer		
THIS DOCUMENT LISTS OBSERVATIONS MADE BY THE FDA REPRESENTATIVE(S) DURING THE INSPECTION OF YOUR FACILITY. THEY ARE INSPECTIONAL OBSERVATIONS; AND DO NOT REPRESENT A FINAL AGENCY DETERMINATION REGARDING YOUR COMPLIANCE. IF YOU HAVE AN OBJECTION REGARDING AN OBSERVATION, OR HAVE IMPLEMENTED, OR PLAN TO IMPLEMENT CORRECTIVE ACTION IN RESPONSE TO AN OBSERVATION, YOU MAY DISCUSS THE OBJECTION OR ACTION WITH THE FDA REPRESENTATIVE(S) DURING THE INSPECTION OR SUBMIT THIS INFORMATION TO FDA AT THE ADDRESS ABOVE. IF YOU HAVE ANY QUESTIONS, PLEASE CONTACT FDA AT THE PHONE NUMBER AND ADDRESS ABOVE. DURING AN INSPECTION OF YOUR FIRM (I) (WE) OBSERVED: PRODUCTION SYSTEM			
1.) Blending of in-specification with out-of-specification (OOS) intermediate batches is performed.			
The batch was manufactured starting on U3/05/2018. QC impurity against a specification of less than or equal to this incident, calls for this batch to be blended with other manufacturing. The OOS batch was blended with in-spec A.) kg of batch blended with blended w	testing of this batch yield %. Deviation investigate batches that have an in-sp ification material as follow kg of batch	on DV/ 1 pecification res ws: to manufacture	% for the 801, written for ult for batch
	g of batch (b) (4)	and ^{(b) (4)} kg o	f batch
The manufactured batches were used in the production of packaged lots of of of the production of packaged lots.			
LABORATORY CONTROL SYSTEM 2.) Laboratory investigations for OOS intermediate batches are not conducted. Specifically, according to SOP BDQC-311 for OOS investigations (effective 07/16/2018), OOS is to be conducted on "all quantitative and qualitative tests ofintermediates" An investigation of an OOS impurity result in the ntermediate, batch for (b) (4) (a) (b) (4) (b) (4) (b) (4) (c) (d) (d) (d) (d) (d) (d) (d) (d) (d) (d			
SEE REVERSE Alan P. Digkally signed by Alan P. Kurtzberg -S DN: c=US, o=US. Government,	PLOYEE(S) NAME AND TITLE (Print of land P. Kurtzberg, Investigator	r Туре)	DATE ISSUED 10/26/2018