TECHNICAL SUPPORT SECTION EFFICACY REVIEW - I Disinfectants Branch

IN_2	2-24-83	OUT 3-1-83
	24-0.	

Reviewed By Dorothy M. Portner Date 3-1-83	
EPA Reg. No. or File Symbol 46506-R	
EPA Petition or EUP No.	-
Date Division Received 2-22-83	-
Type Product Hospital Disinfectant	
Data Accession No(s).	جنب
Produc: Manager 32	
Product Name Bionox No. 1	
Company Name The Bionox Corporation	
Submission Purpose Resubmission of proposed label for determination o	£
the data required to support efficacy	
Type Formulation Liquid concentrate to be mixed with equal parts of	
Active Ingredient(s): %	
Sodium hypochlorite 0.	5

200.0 Introduction

200.1 Use

See attached proposed label.

200.2 Background Information

The submission, received 2-22-83 in response to a directive indicated in the conference of 1-12-83, included a proposed label for representation of this product solely as a hard surface disinfectant. Since the efficacy claims and the patterns of use intended for this product were not precisely delineated at this conference, the data requirements to support product efficacy were deferred until receipt of proposed labeling.

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TECHNICAL SUPPORT SECTION EFFICACY REVIEW - II Disinfectants Branch

EPA Reg. No. or File	Symbol 46506-R	•	
Date Division Receiv	red 2-22-83		•
Data Accession No(s)	•		
Product Manager No	32		
Product Name	Bionox No. 1		
Company Name	The Bionox Corporation		

202.0 Recommendations

202.1 Data Required To Support Efficacy Claims

Based on the efficacy claims and pattern of use indicated on the proposed labeling submitted, the data indicated in (c) of the DIS/TSS-1 enclosure are required. The testing should be conducted for a 10-minute contact time by the AOAC Germicidal Spray Products Test in accordance with the use directions for the spray equipment to be employed with this two component product. If disinfection is intended in less than 10 minutes, as claimed on the center panel of the proposed label, all required data should be developed at the shortest contact time to be claim instead of 10 minutes as specified in the method. A complete description of all models of spray equipment to be marketed, including the directions for use, is required for the file record since this equipment is collateral to the use of this product. The information indicated in the DIS/TSS-3 enclosure is required for reporting of data, including the neutralization procedures employed as indicated in the DIS/TSS-2 enclosure. The data and requested information should be included in the same submission for expediency of the review.

202.2 Labeling

Any revisions required in the proposed labeling will be indicated after the efficacy data have been reviewed.