

1.	PRODUCT AND COMPANY IDENTIFICATION				
(a)	Product Name:	Avioq HIV-1 Microelisa System			
(b)	Product Number:	100384, 100960, 109600			
(c)	Product Description:	The Avioq HIV-1 Microelisa System is an enzyme-linked immunosorbent assay (ELISA) for the qualitative detection of antibodies to Human Immunodeficiency Virus Type 1 (HIV-1) in human specimens collected as serum, plasma, dried blood spots, or oral fluid specimens obtained with Oracure® HIV 1 Oral Specimens			
		with OraSure®.HIV-1 Oral Specimen Collection Device.			
(d)	Supplier Information:	Avioq 104 T.W. Alexander Drive Research Triangle Park, North Carolina 27709 Telephone: 919-314-5535 Fax: 919-314-5536			
(e)	Emergency Phone Number:	Telephone: 919-314-5535			
2.	HAZARDS IDENTIFICATIO	N			
(a)	Classification of the subst	ance or mixture:			
	Kit Component	Description	¹ Hazardous Components of Mixture Present Above Threshold Levels		
	HIV-1 Microelisa Strips	Twelve per holder, each containing ² HIV-1 antigen coated wells, including inactivated, purified viral lysate protein, contained in a foil pack (2 stripholders/pack) with silica gel desiccant.	None		
	Dilsim™ III (120 ml bottle)	Liquid specimen diluent; contains bovine and caprine proteins, salt, surfactant, Chlorophenol Red as specimen addition indicator, Triton x-100 and antimicrobial agents (contains 0.03% [w/v] bromonitrodioxane),	Triton X-100, CAS 9002-93-1: 1.4% Serious Eye Damage/Eye Irritation (Category 2), H319		
	Negative Calibrator Serum (Human) (0.5 ml vial)	Contains ² human serum and goat serum as stabilizer with 0.05% (w/v) bromonitrodioxane as preservative and Fast Green FCF as coloring agent; no detectable antibodies to HBsAg, HIV-1 or HCV.	None		



	HIV-1 Positive Control Serum (Human) (0.5 ml vial)	Inactivated ² human serum containing protein stabilizers. Contains 0.05% (w/v) bromonitrodioxane as preservative and coloring agent; reactive for antibodies to HIV-1 and nonreactive to HBsAg and antibodies to HCV.	None	
	Peroxidase Conjugated Goat Anti-human Immunoglobulins (EnzAbody®) (vial)	Lyophilized with protein stabilizers and FD&C red dye no. 2.	None	
	Conjugate Diluent (120 ml bottle)	Phosphate buffered saline containing protein stabilizers and 0.03% (w/v) bromonitrodioxane as preservative.	None	
	ABTS Substrate Solution (62 ml bottle)	2,2'-azino-di-[3- ethylbenzthiazoline-6-sulfonate], containing hydrogen peroxide.	None	
	Stop Solution (120 ml bottle)	Contains 0.28% sodium fluoride and 0.03% (w/v) bromonitrodioxane as preservative.	None	
	contains less than 1% of a be considered hazardous.	azard Communication Standard (29 CFF) hazardous chemical or 0.1% of a carcing Under GHS, the cutoff level for repropery 1 mutagenicity is ≥0.1%. The cutoff	nogen, the mixture shall not ductive toxicity,	
(b) GHS label elements, precautionary statements:			ments:	
	Kit Component GHS Label Elements, Precautionary Statements:			
	Dilsim™ III (120 ml bottle) Contains: 1.4% Triton X-100			
		Signal Word: Warning Hazard Statements H319 Causes serious eye irritation. Precautionary Statements		



SAFETY DATA SHEET

		<u>, </u>
		P264: Wash hands thoroughly after handling. P280: Wear protective gloves and eye protection.
		Response Precautionary Statements
		P305 + P351 + P338 IF IN EYES: Rinse cautiously with water for
		several minutes. Remove contact lenses, if present and easy to do.
		Continue rinsing.
		P337 + P313 If eye irritation persists: Get medical attention.
(c)	Hazards not otherwise	² According to the OSHA Bloodborne Pathogen Standard (29 CFR
	covered:	1910.1450), all human blood components and certain body fluids must be treated as if known to be infectious for HIV, HBV and other bloodborne pathogens.
		Caution: Handle all Avioq HIV-1 Microelisa System biological
		materials as though capable of transmitting infectious agents. The
		antigen used to coat the microelisa wells and the positive control
		sera have been inactivated; nevertheless, both should be handled
		as though they contain potentially infectious agents. Other
		components prepared from human serum or plasma have been
		tested using FDA-licensed tests and found to be nonreactive for the
		presence of HIV-1 antibody, HTLV-I/II antibody, hepatitis B surface
		antigen (HBsAg) and HCV antibody. However, as no test method
		can offer complete assurance that infectious agents are absent, all
		materials of human origin should be handled as though
		they contain infectious agents.
3.	COMPOSITION	they contain infectious agents.
(a)	Mixtures:	Refer to table of Kit Components in Section 2(a).
4.	FIRST AID MEASURES	(4)
(a)	Route of Exposure	
. ,	In case of skin contact:	Rinse with plenty of water.
	In case of eye contact	See 2(b).
	If inhaled:	Not an expected route of exposure.
	If swallowed:	Rinse mouth. Do not induce vomiting without medical advice. If
		unwell, seek medical attention.
(b)	Symptoms of exposure:	None known.
(c)	Indication of immediate	See 4(a).
	medical attention	
5.	FIRE-FIGHTING MEASURES	
(a)	Extinguishing media:	The components in this kit do not burn. Use extinguishing media that is appropriate for surrounding fire.
(b)	Specific hazards arising	Not considered to be a fire or explosion hazard
	from the product:	
(c)	Special equipment and	None.
	precautions for fire-	
		1
	fighters:	

SDS Number: 410367-02



6.	ACCIDENTAL RELEASE MEASURES			
(a)	Personal precautions, protective equipment, and emergency procedures:	Use suitable personal protective equipment such as gloves, protective clothing or laboratory coat and eye or face protection to avoid skin and eye contact.		
(b)	Methods and materials for containment and cleaning up:	Immediately clean up any spillage of material potentially containing antigen or antibody with a 1:10 dilution of 5% sodium hypochlorite. Dispose of the cleaning material by an acceptable method.		
7.	HANDLING AND STORAG	E		
(a)	Precautions for safe handling:	Avoid contact with skin and eyes. Avoid ingestion and inhalation.		
(b)	Precautions for safe storage:	Store between 2-8°C as indicated in the package insert.		
8.	EXPOSURE CONTROLS/P	ERSONAL PROTECTION		
(a)	Permissible exposure limits:	No data available.		
(b)	Appropriate engineering controls:	General laboratory ventilation is adequate.		
(c)	Personal protective equipment:	Appropriate personal protective equipment should be worn to avoid contact with skin and eyes.		
9.	PHYSICAL AND CHEMICA	L PROPERTIES:		
(a)	Appearance:	Multiple kit components.		
(b)	Odor:	None reported.		
(c)	Odor threshold:	No data available.		
(d)	pH:	No data available.		
(e)	Melting point/freezing point:	No data available.		
(f)	Initial boiling point and boiling range:	No data available.		
(g)	Flash point:	No data available.		
(h)	Evaporation rate:	No data available.		
(i)	Flammability (solid/gas):	No data available.		
(j)	Upper/lower flammability or explosive limits:	No data available.		
(k)	Vapor pressure:	No data available.		
(I)	Vapor density:	No data available.		
(m)	Relative density:	No data available.		
(n)	Solubility:	No data available.		
(o)	Partition coefficient: n-	No data available.		



	octanol/water:		
(p)	Auto-ignition	No data available.	
	temperature:		
(q)	Decomposition temperature:	No data available.	
(r)	Viscosity:	No data available.	
10.	STABILITY AND REACTIVE	тү	
(a)	Reactivity:	No data available.	
(b)	Chemical stability:	No data available.	
(c)	Possibility of hazardous	No data available.	
	reactions:		
(d)	Conditions to avoid:	No data available.	
(e)	Incompatible materials:	No data available.	
(f)	Hazardous	No data available.	
	decomposition		
11.	products: TOXICOLOGICAL INFORM	IATION	
		No data available.	
(a)	Information of likely routes of exposure:	NO data available.	
(b)	Symptoms related to	No data available.	
, ,	the physical, chemical		
	and toxicological		
	characteristics:		
(c)	Delayed and immediate	No data available.	
	effects and also chronic effects form short- and		
	long-term exposure:		
(d)	Numerical measures of	No data available.	
	toxicity:		
(e)	Carcinogenicity:	No component of this product present at levels greater than 0.1%	
42	5001001041 19150094471	are listed by OSHA, IARC, NTP, or CA Prop 65	
12.	ECOLOGICAL INFORMATION		
(a)	Eco-toxicity (aquatic and terrestrial, where	No data available.	
	available):		
(b)	Persistence and degradability:	No data available.	
(c)	Bioaccumulation:	No data available.	
(d)	Mobility in soil:	No data available.	
(e)	Other adverse effects:	This product released into the environment is not expected to have a significant impact.	
13.	DISPOSAL CONSIDERATIONS		
	In accordance with 40 CFR Parts 261.3t this product is not considered to be an EPA		
	Hazardous Waste. Waste generators should always consult with state and local regulations		



	to ensure complete and accurate classification.			
14.	TRANSPORT INFFORMATION			
	Not regulated as a hazardous material.			
15.	REGULATORY INFORMATION			
	SARA 302 Components	No ingredients in this product are subject to the reporting requirements of SARA Title III, Section 302.		
	SARA 313 Components	This product does not contain any ingredients with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.		
	SARA 311/312 Hazards	No SARA Hazards		
	California Prop 65 Components	This product does not contain any chemicals known to the State of California to cause cancer, birth defects, or any other reproductive harm.		
16.	OTHER INFORMATION			
	Kit Component		NFPA Rating	HMIS Rating
	Dilsim™ III (120 ml bottle)		Health Hazard: 1 Fire Hazard: 0 Reactivity Hazard: 0	Health Hazard: 1 Fire Hazard: 0 Physical Hazard: 0