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ANNEX
The following entry 'No 7' is inserted in Annex I to Directive 98/8/EC:

No		n IUPAC			Deadline		Product	_
	Name		e <b>ntifr<del>it</del>a</b> tio	n of	for	date of	type	provisions
		Number		inclusion		n <b>de</b> nclusion	l	•
			active		with			
			substanc	e	Article			
			in the		16(3)			
			biocidal		(except			
			product		for			
			as		products			
			placed		containi	ng		
			on the		more			
			market		than			
					one			
					active			
					substanc	e,		
					for			
					which the			
					tne deadline			
					to			
					comply			
					with			
					Article			
					16(3)			
					shall			
					be the			
					one			
					set out			
					in the			
					last			
					of the			
					inclusion	1		
					decisions	I		
					relating	,		
					to its			
					active			
					substanc	es)		
<b>'</b> 7	carbon	carbon	990 ml/l	1	31	31	14	When
/	dioxide	dioxide	/// IIII/I	November		October	1 I	assessing
	GIONIGO	EC No:			2011	2019		the
		204-696-9		2007		2017		application
		CAS						for
		No:						authorisation
		124-38-9						of a
		12 . 30 )						product
		1	1	1				Product

a For the implementation of the common principles of Annex VI, the content and conclusions of assessment reports are available on the Commission website: http://ec.europa.eu/comm/environment/biocides/index.htm

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	1		,	,	
					accordance
					with
					Article
					5 and
					Annex
					VI,
					Member
					States
					shall
					assess, when
					relevant
					for the
					particular
					product,
					the
					populations
					that
					may be
					exposed
					to the
					product
					and the
					use or
					exposure
					scenarios
					that
					have
					not been
					representatively addressed
					at the
					Community
					level
					risk
					assessment.
					When
					granting
					product
					authorisation,
					Member
					States
					shall
					assess
					the risks
					and
					subsequently
					ensure
					that
					appropriate
					measures

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				are
				taken or
				specific
				conditions
				imposed
				in
				order to
				mitigate
				the
				identified
				risks.
				Product
				authorisation
				can
				only be
				granted
				where
				the
				application
				demonstrates
				that
				risks
				can be
				reduced
				to
				acceptable
				levels.'

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