

Stereospecific Degradation of Diastereomers by Plant Associated Bacteria Influences the Antifungal Performance of Dodemorph*

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Abstract

Morpholine fungicides have certain antibacterial side effect, dodemorph being the most active among them. The diequatorial (*cis*-) form of dodemorph expressed higher antibacterial activity than the axial-equatorial (*trans*-) form, and no synergy in their joint action could be revealed in this respect. Moreover, the partition of diastereomers between cells and medium strictly correlated to their toxicity. Considerable differences were detected among degradation rates in various bacteria, and the *meso*-(*RS*)-diastereomer was deteriorated more intensively, then the *trans*-(*SS* and *RR*)-forms in *Corynebacterium betae*, *Erwinia uredovora* and *Pseudomonas fluorescens*. As a result, the stereospecific degradation of diastereomers changed their ratio in the medium, thus this metabolic step could influence the antifungal performance of dodemorph based preparations against filamentous fungi. It was demonstrated that due to synergic joint action, the fungistatic effect of morpholine derivatives noticeably increased against *Botrytis cinerea* by changing the ratio of diastereomers.

Keywords

Fungicide, Diastereomer, Biodegradation, Bacteria, Fungi, Synergy, Stereoselectivity

1. Introduction

The morpholine derivatives have been introduced against powdery mildews [1]. These compounds inhibit steroid biosynthesis [2] [3] and lipid peroxidases [4]

*Stereospecific degradation of dodemorph by bacteria.

[5], moreover, they can disturb the semipermeability of plasmalemma by selectively coupling to phospholipids [6] and alter the activity of Mg^{2+} -dependent ATP-ases as well [7]. Their use was declined after discovery of benzimidazole [8] [9] and triazole fungicides [10], however, due to rapid development of acquired tolerance to highly specific monosite inhibitors in populations of target microbes, the importance of morpholines is recently increasing. The morpholine fungicides exhibit activity against Gram positive bacteria [2] and some Gram negative species [11] [12] being stereospecific in most of cases [13].

The aim of these studies was to elucidate the importance of stereochemistry of the methyl groups on morpholine ring (**Figure 1**) with special regard to possible consequences in fungicidal performance of dodemorph.

2. Materials and Methods

2.1. Compounds and Microbes Tested

Marketed products, Calixin 75 ec (tridemorph, CASN: 24602-86-6), Corbel 75 ec (fenpropimorph, CASN: 67564-91-4) and Meltatox 40 ec (dodemorph, CASN: 1593-77-7) as well as their active substances were used for testing antimicrobial activities. The analytical standards were gifted by producer (BASF, Ludwigshafen, BRD).

The diastereomers were isolated as follows: Appropriate amount of formulation (4 g of suspected active substance) was added to aqueous sodium carbonate (15% w/v; 100 mL) and the mixture shaken with ethylacetate + cyclohexane (1:1 v/v; 200 mL). The organic phase was removed and treated with active carbon (200 mg) and magnesium sulphate (5 g), then the volume was reduced to 15 mL in rotary evaporator. This mixture was subjected to column chromatography on Kieselgel H, the diastereomers being eluted with dichloroethane + cyclohexanone + ethylacetate + acetone (48 + 50 + 10 + 2 by volume), and were identified by their physicochemical characteristics [10].

The microbes used in toxicity were of the collection of Plant Protection Institute HAS (WDCM824).

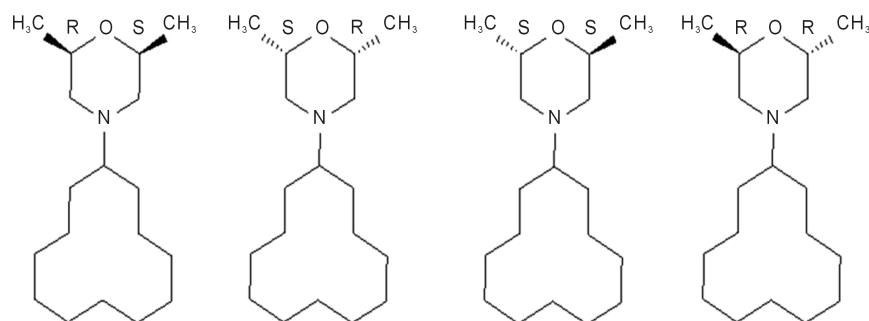


Figure 1. Enantiomers of dodemorph. ACD ChemSketch 2012 was used as graphic tool for drawing and stereochemical labeling. Only one enantiomer (*RS*) exists in the case of *cis*-(diequatorial)-diastereomer (*meso*-form) due to inner symmetry at $N \rightarrow O$ axis in morpholine ring, contrary to *trans*-(axialequatorial)-form, which has two enantiomers (*SS* and *RR*, respectively).

2.2. Toxicity Tests

The bacteria (listed in **Table 1**) were maintained on nutrient agar slants and cultured in nutrient broth containing Bacto peptone, Yeast extract (Difco), glucose and glycerol (10, 1, 3 and 1 g·L⁻¹, respectively) and KH₂PO₄, KCl, MgSO₄, CaCl₂, NaCl and Fe³⁺ citrate (0.55, 0.425, 0.125, 0.125, 0.1 and 0.005 g·L⁻¹, respectively). Hoagland Arnon's micoelement solution was also added (1 mL pro 1 L). Solid media were prepared by incorporating agar-agar (Oxoid; 15 g·L⁻¹) into this solution.

Standardized bacterial suspensions for minimal inhibitory concentrations (MIC), and “degradation” tests were prepared by washing cells from 20 h old culture on solid medium (5 mL in a tube) with sterile liquid medium (5 mL). The optical density of the suspensions was determined (520 nm; Unicam SP 800) and water than added to give a final OD₆₀₀ of 0.4 - 0.5. Such suspensions were prepared for each species tested, and added to wells of multipoint inoculator cooled to 1°C - 4°C.

Minimum inhibitory concentrations for bacteria were determined by adding the appropriate compound in sterile distilled water to nutrient agar (30 mL; 45°C), mixing well and dispensing aliquots of the resultant mixture into each of two Petri dishes. These were inculcated with the standard suspension of the appropriate bacteria using a multipoint inoculator. Dishes were incubated (25°C ± 2°C) and bacterial growth evaluated after 48 and 96 hours on the scale 1 - 4, where 1 represented stimulation, 2 similar to and 3 retardation of growth compared with that on the control and 4 represented no growth. A series of concentration (1000/2ⁿ mg·L⁻¹, n = 0, 1, 2, ..., 21) were tested.

The fungi, *Aspergillus niger* van Tiegh., *Fusarium graminearum* Schwabe, *F. oxysporum* Schlechtendahl, *Chaetomium globosum* Kunze, *Botrytis cinerea* Pers., *Thielaviopsis basicola* (Berk. & Broome) Ferraris, *Rhizoctonia solani* Kühn, *Trametes versicolor* (L.) Lloyd, *Pythium irregularare* Buisman and *Phytophthora cactorum* Pethyb. were maintained and cultured on home prepared potato dextrose agar (PDA) slants amended with mineral salts listed above and sodium nitrate (2 g·L⁻¹). The broth was prepared from peeled potato tubers (200 g) cut into pieces (approx. 1 × 1 × 1 cm) and cooked in 1 litre of distilled water (20 min) then filtered through a Perlon meshwork (No. 500) and the filtrate was then made up to 1 litre with distilled water.

The fungotoxicity of compounds was characterized by ED₅₀ values (mg·L⁻¹), calculated from radial growth inhibition rates expressed as percentage of control grown on poison free medium. The synergistic character of joint action of diastereomers was tested according to either Colby [14] or Sun [15]. The comparative toxicity index (Co.T.I.) was calculated using ED₅₀ values as follows: Co.T.I. = (1/F)/((a/C) + (b/T)), where F, C and T are the ED₅₀ values of the commercial product (F) and single diastereomers (T = *trans*- and C = *cis*-, and a and b their weight ratios in the active ingredient of marketed fungicide.

2.3. Degradation Experiments

Standard suspension of bacteria (5 mL) was added to nutrient broth (150 mL) in 500 mL Erlenmeyer flask and the whole incubated on a rotary shaker at 24°C for 22 hours (*A tumefaciens*, *E. uredovora*, *P. fluorescens* and *X. vesicatoria*) or 44 hours (*C. betae*). The resulted suspensions were centrifuged (10 min, 9000 g), the pellet washed with sterile medium (without peptone and yeast extract), and the residue weighed. All operations were performed at 0°C - 1°C, and the medium was adjusted to pH 6.8 prior use. A portion of the cells (500 mg) was added to the medium (25 mL, without peptone and yeast extract) to give the working suspension used for studies of partition coefficient and degradation.

The acetonous stock solution of the appropriate compound (500 µg) as added to the working suspension, mixed well, and the suspension incubated on a rotary shaker at 27°C ± 1°C for 5 hours. The suspension was then cooled melting ice bath to 0°C and centrifuged (10 min, 9000 g) at 0°C - 1°C. The supernatant and residual bacteria were analyzed separately.

2.4. Analytical Techniques

Silufol UV₂₅₄ plates (Cavalier, Votice, Bohemia) with benzene + ethylacetate (78 + 22 v/v) as mobile phase were used for purity test, where diastereomers were visualized with iodine vapor as well, the minimum quantity detectable being 0.1 µg (10⁻⁶ mM).

Packard series 7400 instrument equipped with a flame ionization detector, and 180 × 2 mm I.O. glass column packed with 3% OV-1 + 1.5% OV-225 on Gaschrom Q (80 - 100 mesh) were used for separation of diastereomers. The detector and injector temperatures were 235°C. The column temperature was changed at 10°C min⁻¹ from 150°C to 220°C after 4 min initial hold and 9 min final hold. 5 × 10⁻⁶ g of compound were injected in 5 microlitre, final resolution power 3 × 10⁻⁹ as the final point of the calibration curve.

For determination of the quantity of test compounds in medium of degradation experiments 0.5 mL acetone and 2 mL saturated sodium sulphate containing 5% v/w sodium carbonate was added to the supernatant, which was then extracted with 1.0 mL of cyclohexanone + ethylacetate + toluene (5 + 1 + 5 v/v) and the organic phase used for GC analysis immediately. The degradation rate was calculated from the analytical data as a difference of test substances applied and that in the supernatant and in the bacteria after 5 hours incubation.

The bacterial residue was suspended in distilled water (0.5 mL), then acetone (0.5 mL) added and the mixture kept in boiling water for 3 min. Distilled water saturated with sodium chloride (4 mL) were than added, and the resulting suspension extracted as above. The organic phase was subsequently analyzed by GC. The distribution of test chemicals between bacterial cells and medium was expressed as K;

K = [concentration (microg/g) in the cell]/[concentration (microg/mL) in the medium]

and K was a measure of uptake of the investigated substance into the cells.

2.5. Data Treatment

All experiments were carried out at least in triplicates. Student's *t* was used to evaluate significance of differences between average values, and Fisher's test was applied to reveal impact of alterations in structure of test compounds at P = 5% level.

3. Results and Discussion

3.1. Toxicity

Meltatox proved to be most effective against bacteria surpassing Calixin or Corbel (**Table 1**). In general, the Gram positive bacteria were essentially more sensitive to morpholine fungicides than Gram negatives. Although, no data are available for the intrinsic activities of formulants, seemingly all test microbes tolerated well the carriers used for preparation as only at high concentrations were revealed minuscule differences between commercial products and their active ingredients. The diastereomers had different toxicities (**Table 1**). The most effective *cis*-form seems to be responsible for the toxicity of formulated product, Meltatox 40 ec, *Agrobacterium*, *Erwinia* and *Pseudomonas spp*. The *cis*-form is also more toxic to *Xanthomonas* species, while Gram positives exhibited differential sensitivity.

Differential sensitivity was also demonstrated among fungal species investigated *in vitro* (**Table 2**). Calixin 75ec was most toxic to the majority of fungi, while Meltatox 40ec and the constituent dodemorph diastereomers proved to be the least toxic. Although, in several cases significant difference of activity could not be revealed between diastereomers, their mixtures always exhibited higher toxicity. The synergy varied within large limits, and it was outstanding in the case of *B. cinerea*. The stereoselective antifungal activity is not a proper character of dodemorph as it was similarly elucidated in the case of tridemorph diastereomers as well.

3.2. Uptake of Diastereomers into Bacterial Cells

Uptake of diastereomers into bacterial cells was very different with the representatives of the five genera tested and seems to be proportional to their sensitivity (**Table 3**). The *trans*-form was accumulated much more than the *cis*-form and the ratio varied a little. This enrichment with a particular diastereomer was inversely correlated with lipophilicity over a range of pH. The *cis*-form is 2 - 4 times more lipophilic than the *trans* one. Cells of these bacteria are surrounded with a hydrophilic capsule having acidic character. This capsule is clearly the first barrier to permeation of the molecules into the cells. Perhaps this is the cause of the inverse correlation between the lipophilicity of dodemorph isomers and their accumulation in the cells. The effect of cell wall and mucous capsule can be possibly influenced by the change of the pH which depending on the

Table 1. Growth inhibitory effect of morpholine derivatives towards bacteria.

Bacteria	Minimum inhibitory concentrations (mg·L ⁻¹)					
	Dodemorph		Tridemorph		Fenpropimorph-	Strepto. mycin
	cis-	trans-	cis-	trans-		
<i>Agrobacterium</i>						
<i>A. radiobacter</i> K-84	125 - 250	>1000	250 - 500	250 - 500	>1000	>1000
<i>A. tumefaciens</i> C-58	62 - 125	>1000	125 - 250	250 - 500	>1000	62 - 125
<i>A. tumefaciens</i> O	125 - 250	>1000	125 - 250	250 - 500	>1000	>1000
<i>Bradyrhizobium</i>						
<i>B. japonicum</i> 27	125 - 250	>1000	31 - 62	62 - 125	>1000	>1000
<i>Rhizobium</i>						
<i>R. trifolii</i> 28	250 - 500	>1000	500 - 1000	500 - 1000	>1000	>1000
<i>Erwinia</i>						
<i>E. atroseptica</i>	125 - 250	>1000	31 - 62	31 - 125	>1000	125 - 250
<i>E. carotovora</i>	>1000	>1000	31 - 62	15 - 31	>1000	125 - 250
<i>E. chrysanthemi</i> B8	500 - 1000	>1000	500 - 1000	500 - 1000	>1000	1 - 2
<i>E. herbicola</i> D-5	62 - 125	>1000	125 - 250	125 - 250	>1000	31 - 62
<i>E. uredovora</i> 3vfr	500 - 1000	>1000	250 - 500	500 - 1000	>1000	8 - 16
<i>Pseudomonas</i>						
<i>P. fluorescens</i> K-20	500 - 1000	>1000	500 - 1000	500 - 1000	>1000	16 - 31
<i>P. lachrymans</i> 12	>1000	>1000	125 - 250	250 - 500	>1000	1 - 2
<i>P. phaseolicola</i> 14	>1000	>1000	125 - 250	16 - 31	>1000	>1000
<i>Xanthomonas</i>						
<i>X. alfalfae</i> KK-1	250 - 500	500 - 1000	125 - 250	250 - 500	>1000	4 - 8
<i>X. campestris</i> 2D510	125 - 250	16 - 31	62 - 125	125 - 250	500 - 1000	16 - 31
<i>X. pelargonii</i> 58	8 - 16	8 - 16	31 - 62	125 - 250	62 - 15	16 - 31
<i>X. vesicatoria</i> 53	125 - 250	250 - 500	125 - 250	125 - 250	>1000	16 - 31
<i>Corynebacterium</i>						
<i>C. betaee</i> 101	31 - 62	8 - 16	31 - 62	31 - 62	31 - 62	n.t.
<i>C. fascians</i> 27	1 - 2	0.01 - 4	16 - 31	16 - 31	31 - 62	62 - 125
<i>C. flaccumfaciens</i> 8	62 - 125	62 - 125	16 - 31	31 - 62	31 - 62	16 - 31
<i>C. michiganense</i> 9	4 - 8	0.1 - 0.2	2 - 4	8 - 16	31 - 62	1 - 2
<i>C. nebraskense</i> 101	4 - 9	0.5 - 1	2 - 4	16 - 31	31 - 62	125 - 250
<i>C. oortii</i> 11	8 - 16	16 - 31	8 - 16	31 - 62	31 - 62	16 - 31

Code numbers of *Agrobacterium*, *Bradyrhizobium*, *Rhizobium*, *Erwinia*, *Pseudomonas*, *Xanthomonas* and *Corynebacterium* strains were given by identifiers. n.t. = not tested.

Table 2. Inhibitory effect of morpholine fungicides and their diastereomers on the radial growth of fungi.

Fungi	Meltatox 40 ec	Dodemorph		Co. T.I.	Calixin 75 ec	Tridemorph		Co. T.I.	Corbel 75 ec
		cis	trans			cis	trans		
<i>P. irregularare</i>	32.7	60.5	21.2	1.1	27.0	34.9	60.7	1.7	58.2
<i>P. cactorum</i>	178.7	890.1	89.0	1.1	3.6	23.1	41.3	8.5	26.5
<i>A. niger</i>	151.4	572.0	223.6	2.3	3.8	8.6	8.2	2.2	8.7
<i>F. graminear.</i>	38.0	77.4	70.6	2.0	14.5	39.3	29.0	2.3	9.7
<i>F. oxysporum</i>	43.7	112.1	122.9	2.7	6.8	35.8	32.6	5.0	27.5
<i>C. globosum</i>	7.94	70.70	50.10	7.65	1.86	2.18	5.83	1.79	0.76
<i>B. cinerea</i>	0.33	7.8	14.8	28.9	0.31	1.5	2.0	5.5	0.2
<i>T. basicola</i>	17.4	60.2	72.3	3.7	1.1	3.6	5.4	4.1	1.8
<i>R. solani</i>	3.6	7.8	4.5	1.6	1.0	3.2	4.5	3.9	0.15
<i>T. versicolor</i>	5.0	7.8	30.9	2.2	0.64	0.90	3.0	2.3	0.19

The body of the table comprises ED₅₀ values (mg·L⁻¹) and comparative toxicity indices characterizing the joint action of diastereomers. Corbel 75ec contains only cis-fenpropimorph. Example for calculating Co. T.I. = (1/0.33)/((0.6/7.8) + (0.4/14.8)) = 28.9 in the case of *B. cinerea*. The joint action is synergic when Co. T.I. > 1.0, and antagonistic when Co. T.I. < 1.0, while the effect is considered to be additive when Co. T.I. = 1 ± 0.05.

Table 3. Uptake and degradation of dodemorph diastereomes in bacterial suspension.

Parameters	Isomers	Bacteria ^a					LSD _{0.05}
		A	E	P	X	C	
Measured quantity (μg) in							
Cells	cis-D	66.5	24.8	21.5	66.9	156.9	4.94
	trans-D	91.8	46.1	77.5	96.5	174.6	
	c/t	0.72	0.54	0.28	0.69	0.90	
Medium	cis-D	45.6	207.4	187.2	94.2	36.1	4.94
	trans-D	22.4	159.1	116.5	47.1	16.2	
	c/t	2.04	1.3	1.61	2.00	2.23	
Distribution (K = [cell]/[medium])							
	cis-D	29.2	2.4	2.3	14.2	87.0	
	trans-D	81.6	5.8	12.8	40.9	216.1	1.49
	c/t	0.36	0.41	0.18	0.35	0.40	
Degradation (μg/g cells/hour)							
	cis-D	162.9	42.7	66.3	113.9	81.9	21.9
	trans-D	110.7	19.8	34.1	81.8	34.2	
	c/t	1.47	2.16	1.94	1.39	2.39	
Ratio of ionized diastereomers (%) ^b							
Actual pH		4.9	4.7	4.1	5.5	4.5	
	cis-D	98.32	98.84	99.68	93.5	99.31	
	trans-D	96.84	97.49	99.35	86.01	98.40	

^a*A. tumefaciens* C58, *E. carotovora*, *P. fluorescens*, *X. vesicatoria* and *C. betae* were used for experiments. ^bThe medium was acidified during the incubation at species dependent manner. The pH of the bacterium suspensions was 6.8 at the start of incubation, and the degree of ionization was 50 and 28.9 % of cis- and trans-diastereomers, respectively.

species drops during the incubation from the original 6.8 to 4.1 - 5.5. We can suppose the existence of pH gradient (changing during incubation) via medium/outside of mucous capsule to inner side of plasmalemma, which can change the degree of ionization of capsule-cell wall and both diastereomers. In the last case the change of pH in the medium moves the ratio of cationic form from about 25% to 80% - 99% in dependence of the species and diastereomeric form concerned (pK_b values for *cis*- and *trans*-forms are 6.66 and 6.29, respectively). As the difference between accumulation of *cis*- and *trans*-forms much higher than the difference between their ionization in each proper case, we can suppose, that the selective "accumulation" observed was not due to the fact, that the increased ionization upraises the possibility of bounding a cation, so the inverse "lipophilicity" correlation is not an artifact.

3.3. Degradation

The quantity of both diastereomers was diminished by bacteria in the incubation system during a 5 hours incubation. This phenomenon we could detect only in case of intact cells. For this we suggest that it is the result of the transformation, character of what we did not investigate, because of the instability of the compounds of degradation, however, the original molecules we could resolve with high degree. This made possible to extract information from the analytical data about degradation as the diminishing of quantity can be calculated, that means, the lack of the total quantity of the substance is an indirect characteristic of the transformation activity in bacteria. Degradation rates in absence of stereospecificity should correspond to the original ratio of diastereomers.

Differences in accumulation of individual isomers per se cause alteration in ratio (*cis*[RS]/*trans*[RR]/*trans*[SS]) of (3 + 3 + 2 + 2) or (*cis*[R,S] versus *trans*[RR, SS]) of 3:2. The ratios calculated from the measured values did not correspond to the original one that was altered to the side of lower values speaking about the more rapid transformation of the more active *cis*(R,S)-form (Table 3). This suggests that the degradation in bacteria means detoxification. The transformation systems of investigated bacterial species distinguished the diastereomers in various degrees, unfortunately, we could not determine the possible change of ratios of RR/SS enantiomers of *trans*-form. From measured values after 5 hours incubation it can be supposed, that one of them is more rapidly transformed during 5 hrs of incubation because of the absence of transformation rate significantly more than 50% of transformation.

The *cis*-isomer was significantly more active than the *trans* one ($t_{C,T} = 7.72$, $p < 0.05$), nevertheless, their ratio in commercial product (3C + 2T) resulted synergic effect in the case of numerous fungi (Table 2) that can influence significantly the fungicidal performance of marketed product.

The biodegradation not only diminishes the quantity of dodemorph applied, but this stereoselective metabolic process also leads to changes in the ratio of synergically interacting diastereomers. Two optimal mixtures with dominating either *cis*- or *trans*-diastereomers were found as evaluated with *B. cinerea* (Figure 2).

The growth inhibitory effect of 9:1 and 3:7 *cis/trans* mixtures did not differ significantly (47.1 ± 2.4 and $45.6\% \pm 3.2\%$, respectively, $t = 0.74$, $p > 0.1$), and the increase was economically acceptable in both cases. Thus, the most rapid degradation of *cis*-form as compared to *trans*-form does not result decreased efficacy of treatment, as the alteration of their ratio of 3:2 to domination of *trans*-form leads to increase of fungicidal activity, that means, the stereoselective deterioration by bacteria may be presumed as a special type of biactivation. Similar interaction could be revealed between *N*-alkyl derivatives as well, and the mixtures of corresponding dodemorph and tridemorph diastereomers also formed synergic compositions.

3.4. Future Prospects

The development of molecular design opened possibility to reinvestigate the structure/activity relationships of morpholine derivatives, which have special place among systemic fungicides due to lack of acquired tolerance in populations of target phytopathogens. The main targets in filamentous fungi are in steroid metabolism, however, their antimicrobial activity relates also to inhibition of

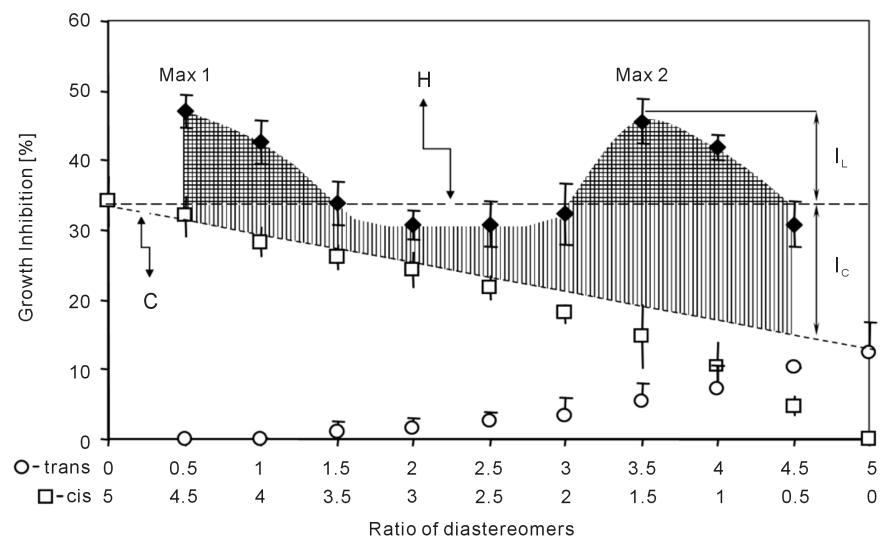


Figure 2. Evaluation of joint action of dodemorph diastereomers against benomyl-tolerant *Botrytis cinerea*. The experimental model of Horsfall (1945) was applied at 5 mg L^{-1} of active ingredient in the PDA. Lines C and H mark additive effect calculated according to Colby [14] and requested minimum increase in efficacy of mixture requested by Lemkin's case [16], respectively. I_L is the maximum increase in efficacy of mixtures evaluated by Horsfall's model [17] and accepted by US Patent Office as economically approved synergy, while I_C is the part of efficacy over additive that does not add significant economic benefit when the two compounds applied in mixture ($t_{C10, C7 + T3} = 0.15$, $t_{C5, C3 + T2} = 1.57$, $t_{C10, C5 + T5} = 1.36$, $t_{C3 + T2} = 0.65$, $t_{C10, C1 + T9} = 1.36 < t_{0.1} = 2.35$), according to decision of Worley *et al.* [16]. Example for evaluation expected activity of Meltax 40 ec (3 *cis*+2 *trans* diastereomers) at 5 mg L^{-1} by Colby's model [14]: The expected activity = C + T - (C × T/100) = $24.3 + 1.5 - ((24.3 \times 1.5)/100) = 25.4\%$, but the measured one was 30.9% that is 5.5% more than the expected. However, this synergic increase in efficacy does not result economic benefit because the *cis*-form resulted 34.2% inhibition when applied alone.

various other target sites in metabolism and cell membrane, which can explain their antibacterial activity. Moreover, tridemorph applied in combination with P₄₅₀ inducer changed the character of host/parasite relationship converting the symptomless sensitive state of plant cells to hypersensitive [7] in the case obligate endoparasites. The mechanisms of cell/cell interaction and nutrition of arc-heas residing in plants are white fields, but these pathogens might be target organisms to morpholine derivatives as phospholipids selectively interact with these compounds [6]. The special advantage of this group is the lack of their accumulation in the environment or in the food chain due to rapid biodegradation of morpholine derivatives.

4. Conclusion

Interaction between dodemorph diastereomers against bacteria could not be revealed, while the synergic joint action essentially influences the fungitoxic action of the formulated product (Meltatox 40ec) containing their mixture. The stereoselective biodegradation of diastereomers in the case of these compounds is a special type of activation that significantly improves the antifungal performance of dodemorph.

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