QSAR model for human intestinal absorption (v1.0)



ProtoADME

ProtoADME is a computational (in silico) tool focused on the prediction of endpoints related with the ADME (Absorption, Distribution, Metabolism and Excretion) of chemical substances.

Endpoint

Toxicokinetic: human intestinal absorption

The human intestinal absorption of an oral drug is the essential prerequisite for its apparent efficacy. The close relationship oral bioavailability and intestinal absorption has also been proven and HIA can be seen an alternative indicator for oral bioavailability to some extent.

Metrics

Training set

Experimental values	QSAR predictions		
	Negative	Positive	
Negative	130	17	
Positive	25	755	

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Experimental values	QSAR predictions					
	Negative	Positive				
Negative	39	11				
Positive	10	252				

Parameters	Training	Validation	
Accuracy	0.95	0.93	
Sensitivity / recall	0.97	0.96	
Specificity	0.88	0.78	
Precision	0.98	0.96	
Negative predictive value	0.84	0.80	
F-score	0.97	0.96	
Matthews Correlation Coefficient	0.83	0.75	
Critical Success Index	0.95	0.92	
Area under the ROC	0.93	0.87	

ProtoADME is part of



ProtoPRED platform allows the easy, fast and user-friendly prediction of different properties of chemical compounds, using proprietary (Q)SAR models.



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