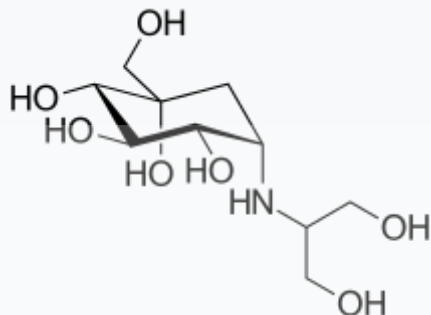


# Voglibose

## Voglibose



## Clinical data

[AHFS/Drugs.com](#)

[International Drug Names](#)

Voglibose 0.2mg / 0.3mg Mouth Dissolving Tablets

[ATC code](#)

[A10BF03 \(WHO\)](#)

## Identifiers

[IUPAC name](#)[\[show\]](#)

[CAS Number](#)

[83480-29-9](#) ✓

[PubChem](#) CID

[444020](#)

[DrugBank](#)

[DB04878](#) ✓

[ChemSpider](#)

[392046](#) ✓

[UNII](#)

[S77P977AG8](#)

[KEGG](#)

[D01665](#) ✓

[ChEMBL](#)

[CHEMBL476960](#) ✓

## Chemical and physical data

[Formula](#)

$C_{10}H_{21}NO_7$

[Molar mass](#)

267.28 g/mol

[3D model \(JSmol\)](#)

[Interactive image](#)

[SMILES](#)[\[show\]](#)

Vobrryl<sup>®</sup> 0.2 / 0.3 Mouth Dissolving Tablet

[InChI\[show\]](#)

[\(verify\)](#)

**Voglibose** ([INN](#) and [USAN](#), trade name **Vogliib**, marketed by Mascot Health Series) is an [alpha-glucosidase inhibitor](#) used for lowering post-prandial blood glucose levels in people with [diabetes mellitus](#). Voglibose delays the absorption of glucose thereby reducing the risk of macrovascular complications. Voglibose is a research product of [Takeda Pharmaceutical Company](#), Japan's largest pharmaceutical company. Voglibose was first launched in 1994, under the trade name BASEN, to improve postprandial hyperglycemia in [diabetes mellitus](#).<sup>[1]</sup>

[Postprandial hyperglycemia](#) (PPHG) is primarily due to first phase insulin secretion. Alpha glucosidase inhibitors delay glucose absorption at the intestine level and thereby prevent sudden surge of glucose after a meal.

There are three drugs which belong to this class, [acarbose](#), [miglitol](#) and voglibose, of which voglibose is the newest.

## References<sup>^</sup>

[https://www.takeda.com/news/2008/20080526\\_3621.html](https://www.takeda.com/news/2008/20080526_3621.html)

