

Simple.  
Every day.

 **Trajenta**<sup>®</sup>  
(linagliptin) 5mg tablets

 **Jentadueto**<sup>®</sup>  
(linagliptin/metformin HCl)

TRAJENTA®



One dose,  
Once daily.<sup>1</sup>



**NO DOSE REDUCTION**  
regardless of renal function<sup>1</sup>



**PROVEN EFFICACY**  
at any stage of renal function<sup>1,2</sup>



Suitable for  
**A BROAD RANGE**  
of T2D patients<sup>1</sup>

1. TRAJENTA® Summary of Product Characteristics. June 2017. 2. McGill JB, et al. Diabetes Care. 2013;36:237-44.



# TRAJENTA®: The only approved\* DPP4i that does not require dose reduction based on renal function<sup>1-5</sup>

Required DPP4i dose with declining renal function, as defined by SmPC<sup>1-5</sup>



	Normal	Mild RI	Moderate RI	Severe RI	End-stage renal disease
<b>5 mg OD</b>					
	100 mg OD	100 mg OD	100 mg OD	50 mg OD	25 mg OD
		GFR ≥60 to 90 mL/min	GFR ≥45 to 60 mL/min	GFR ≥30 to 45 mL/min	GFR ≥15 to 30 mL/min
	5 mg OD	5 mg OD	2.5 mg OD	2.5 mg OD	Not recommended
		CrCl >50 to ≤80 mL/min	CrCl ≥30 to ≤50 mL/min	CrCl > 15 to <30 mL/min	CrCl < 15 mL/min
	50 mg BD	50 mg BD	50 mg OD	50 mg OD	50 mg OD
		CrCl ≥50 mL/min	CrCl ≥30 to <50 mL/min	CrCl > 15 to <30 mL/min	ESRD on haemodialysis: use with caution
	25 mg OD	25 mg OD	12.5 mg OD	6.25 mg OD	6.25 mg OD
		CrCl >50 to ≤80 mL/min	CrCl ≥30 to ≤50 mL/min	CrCl > 15 to <30 mL/min	Limited experience in renal dialysis. Not studied in peritoneal dialysis

When renal function declines, no need for DPP4i dose adjustment with TRAJENTA®<sup>1</sup>

1. TRAJENTA® Summary of Product Characteristics. June 2017. 2. Januvia® Summary of Product Characteristics. December 2017. 3. Onglyza® Summary of Product Characteristics. August 2017. 4. Galvus® Summary of Product Characteristics. April 2017. 5. Vipidia® Summary of Product Characteristics. January 2015.

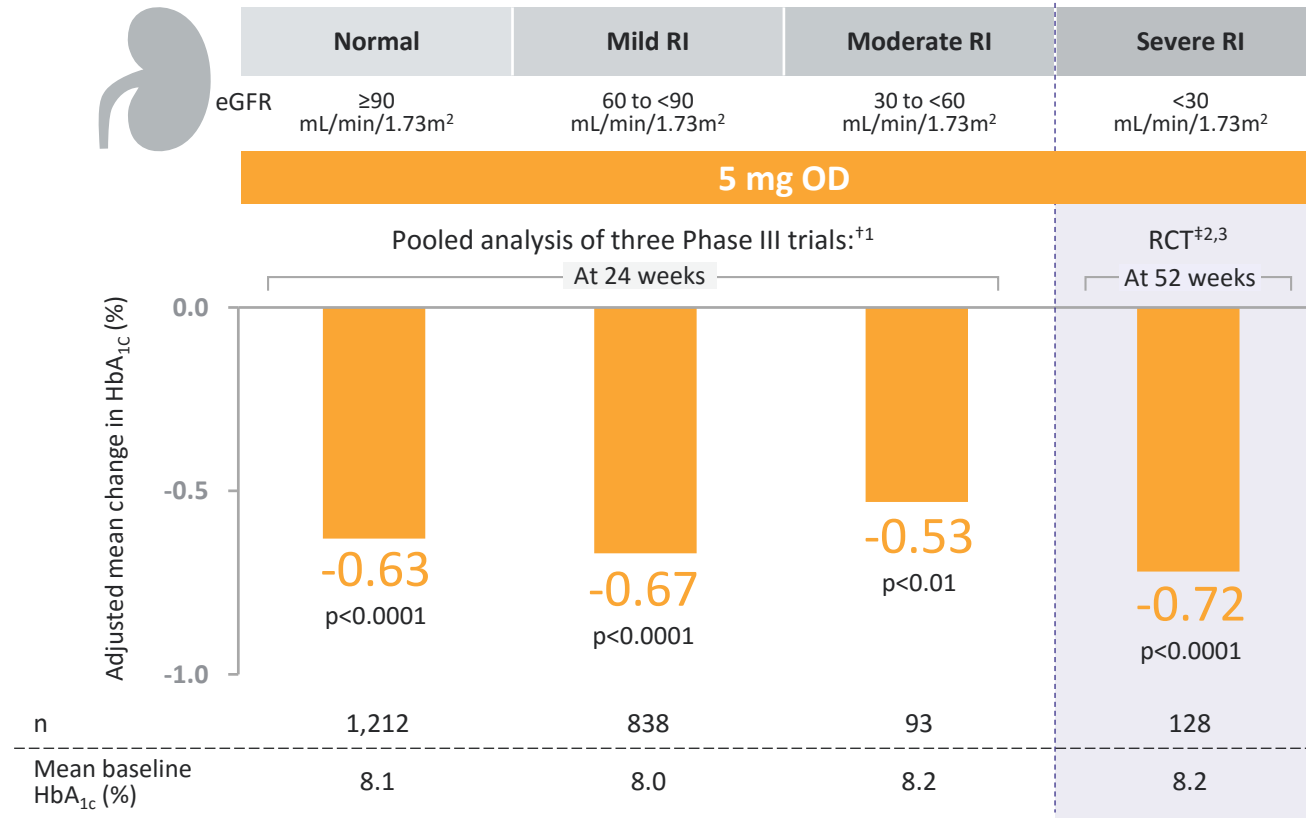
\*Information is derived from the Summary of Product Characteristics (SMPC) for TRAJENTA®/JENTADUETO® in the EU. It is not country-specific and may vary from the approved label in the country where you are located. Please refer to your local Prescribing Information for full details.

BD: Bi-daily; CKD: Chronic kidney disease; CrCl: Creatinine clearance; OD: Once daily; RI: Renal impairment.



# TRAJENTA®: Proven efficacy with the same dose regardless of renal function<sup>1-3</sup>

Adjusted mean HbA<sub>1c</sub> change vs placebo from baseline by degree of renal impairment (RI)\*



1. Groop PH, et al. Diabetes Obes Metab. 2014;16:560-8. 2. McGill JB, et al. Diabetes Care. 2013;36:237-44. 3. TRAJENTA® Summary of Product Characteristics. June 2017.

\*A small proportion of patients in these studies were receiving treatment combinations that fall outside of the licensed indications for TRAJENTA® (linagliptin).

<sup>†</sup>Prespecified subgroup analysis on pooled data from three pivotal Phase III, randomised, placebo-controlled trials: treatment in monotherapy, add-on to metformin, and add-on to metformin plus sulphonylurea. P values for between-group difference (versus placebo). Model includes continuous baseline HbA<sub>1c</sub>, baseline body mass index (category), washout period, treatment, study, age group, gender, time since diagnosis of diabetes, race, renal function (MDRD), and treatment by renal function (MDRD).

<sup>‡</sup>1-year, randomised, double-blind, placebo-controlled study: treatment added to existing background therapy. Data based on analysis using LOCF (last observation carried forward).

eGFR (mL/min/1.73m<sup>2</sup>) = estimated glomerular filtration rate calculated as  $175 \times (\text{Scr})^{-1.154} \times (\text{Age})^{-0.203} \times (0.742 \text{ if female}) \times (1.212 \text{ if Black})$ ; OD : Once daily; RCT: Randomised controlled trial; Scr: Serum creatinine.



# TRAJENTA®: One dose, once daily for your type 2 diabetes patients\*<sup>1</sup>



## Independent of:



Renal  
function



Hepatic  
function



Age



BMI



Background  
T2D therapy



Disease  
duration



Ethnicity

1. TRAJENTA® Summary of Product Characteristics. June 2017.

\*Indicated for use in adult patients. TRAJENTA® is contraindicated in those with hypersensitivity to any of the active substances or excipients, is not licensed for paediatric use and should not be used in pregnant women.

BMI: Body mass index.

