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Schweizerische Gesellschaft für
Endokrinologie und Diabetologie - SGED



Diabetes Epidemiologie

1/12
people with
DIABETES



1 healthcare

in 9
IS SPENT ON DIABETES

In 2014 diabetes expenditure reached US\$612 billion



IDF DIABETES ATLAS Sixth edition



SANOFI DIABETES



ACCUCHEK



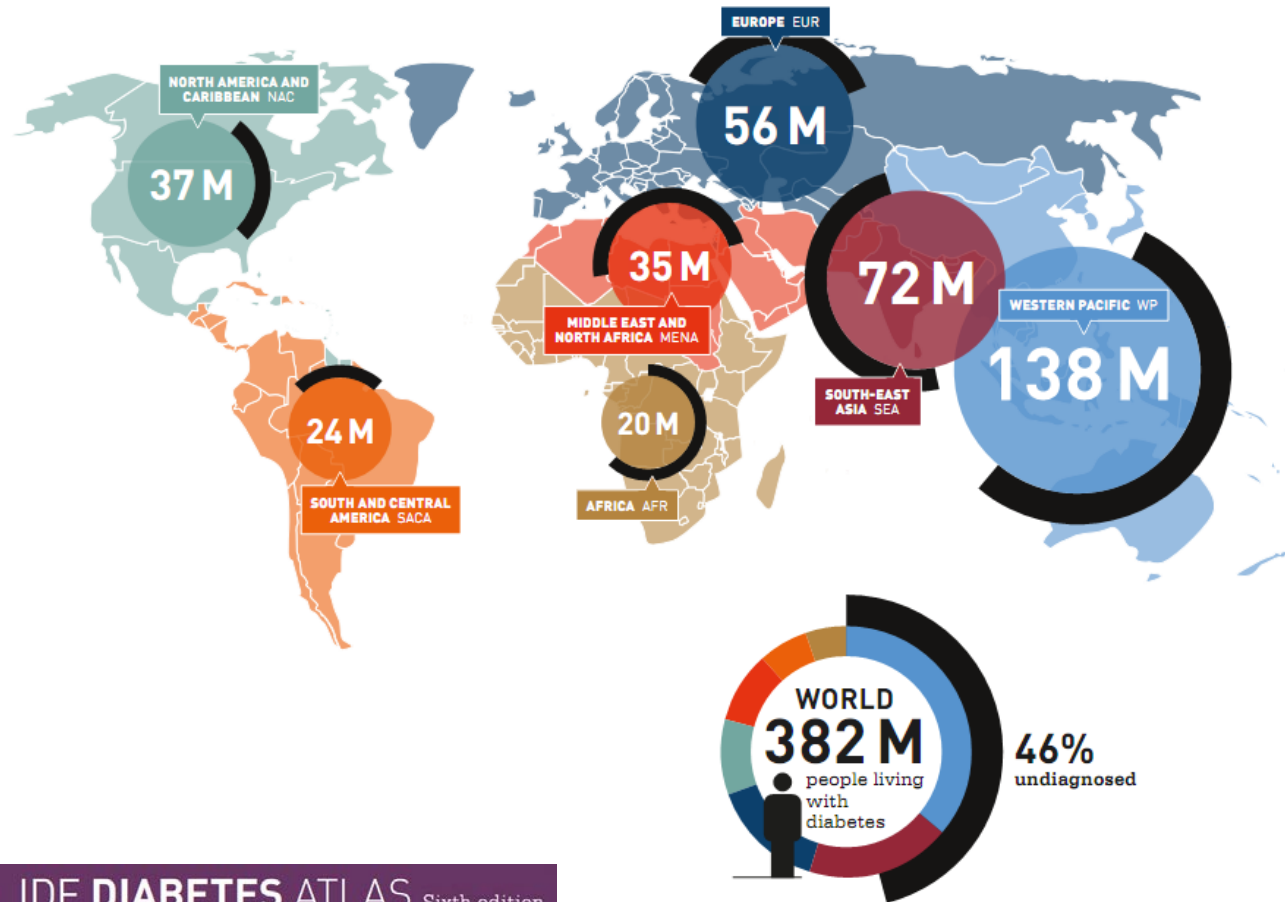
NOVARTIS
PHARMACEUTICALS

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Diabetes Epidemiologie

Number of people with diabetes by IDF Region, 2013



International Diabetes Federation **IDF DIABETES ATLAS** Sixth edition



SANOFI DIABETES



ACCU-CHEK



NOVARTIS
PHARMACEUTICALS

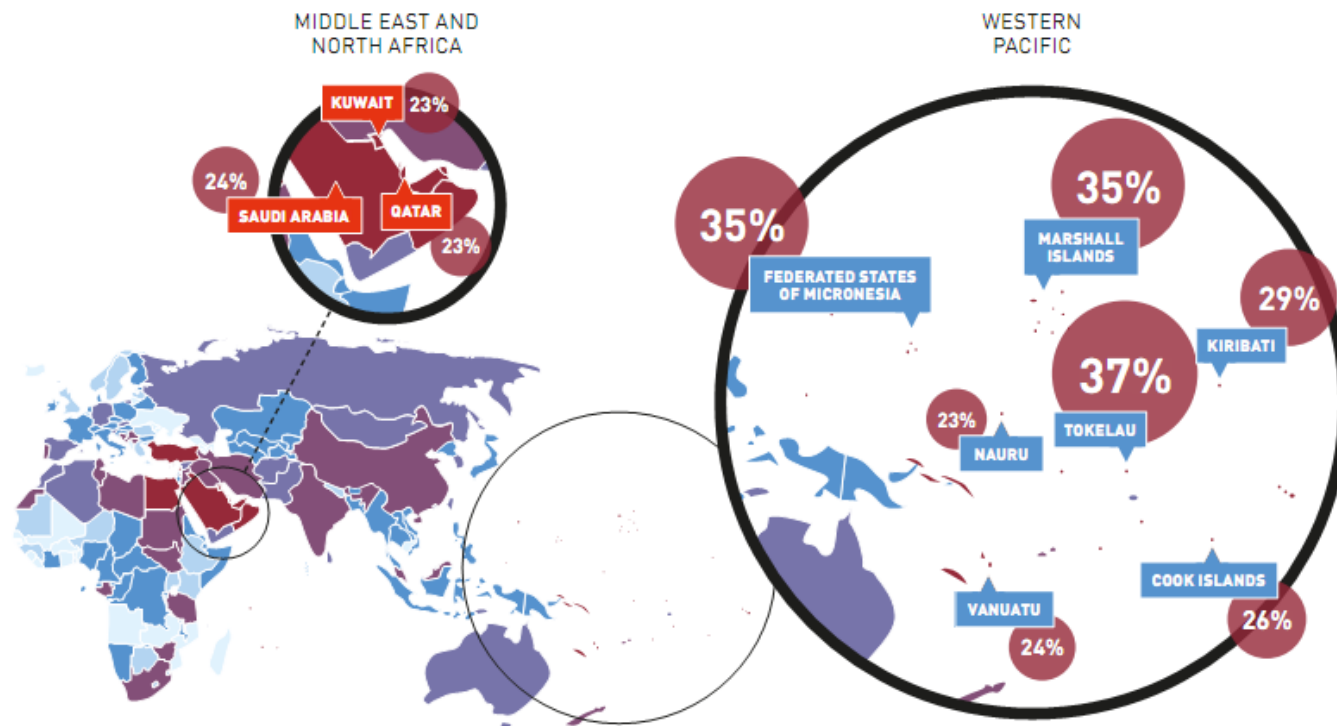
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Diabetes Epidemiologie

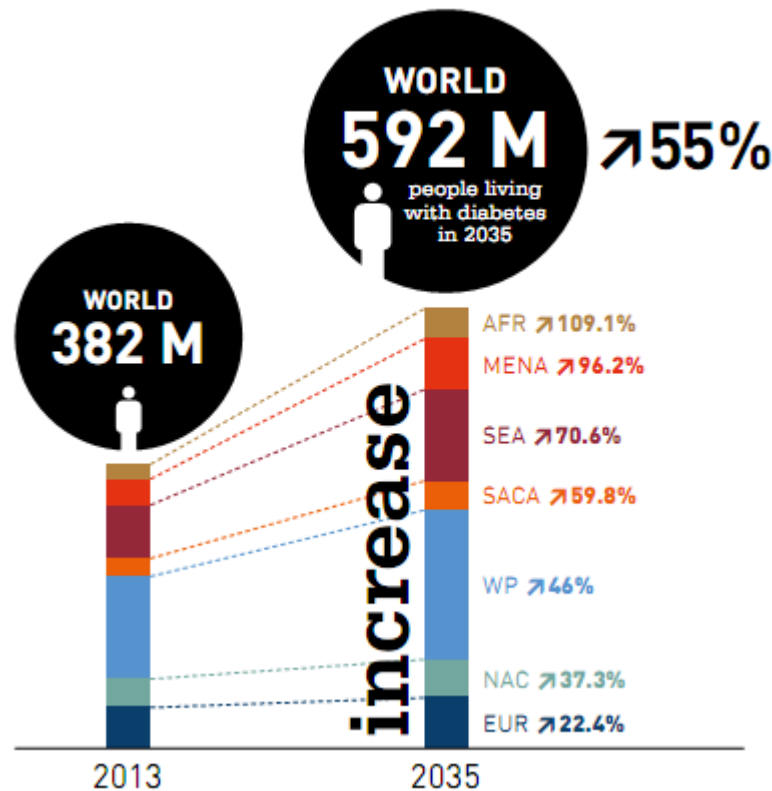
Top 10 countries/territories for prevalence* (% of diabetes (20-79 years), 2013)

* comparative prevalence



International Diabetes Federation **IDF DIABETES ATLAS** Sixth edition

Diabetes Epidemiologie



382 Millionen Personen sind Diabetiker

Bis 2035 werden es 592 Millionen sein



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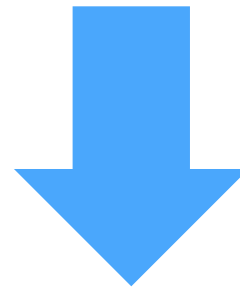


Herausforderungen für den Hausarzt bei Behandlung von Typ 2 DM

- viele Patienten
- viele Komplikationen
- viele (neue!) Medikamente
- viel Geld (vor allem für neue Medikamente)
- intensives Marketing, aber wenig gute Endpunkte



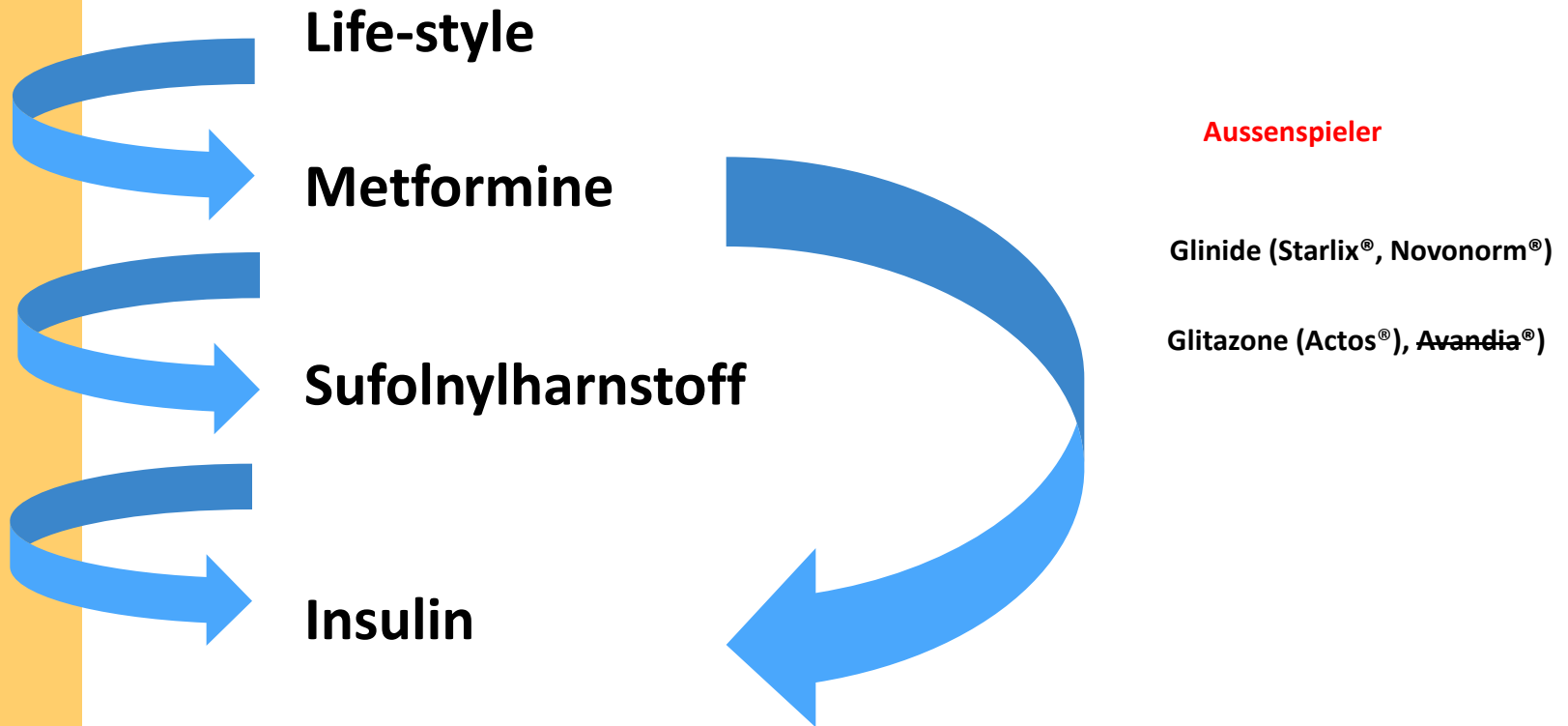
European Society for the Study of Diabetes (EASD) / American Diabetes Association (ADA) Position Statement 2012



**Management of hyperglycaemia in type 2 diabetes, 2015:
a patient-centred approach. Update to a Position Statement
of the American Diabetes Association and the European
Association for the Study of Diabetes**



« klassischer Weg »



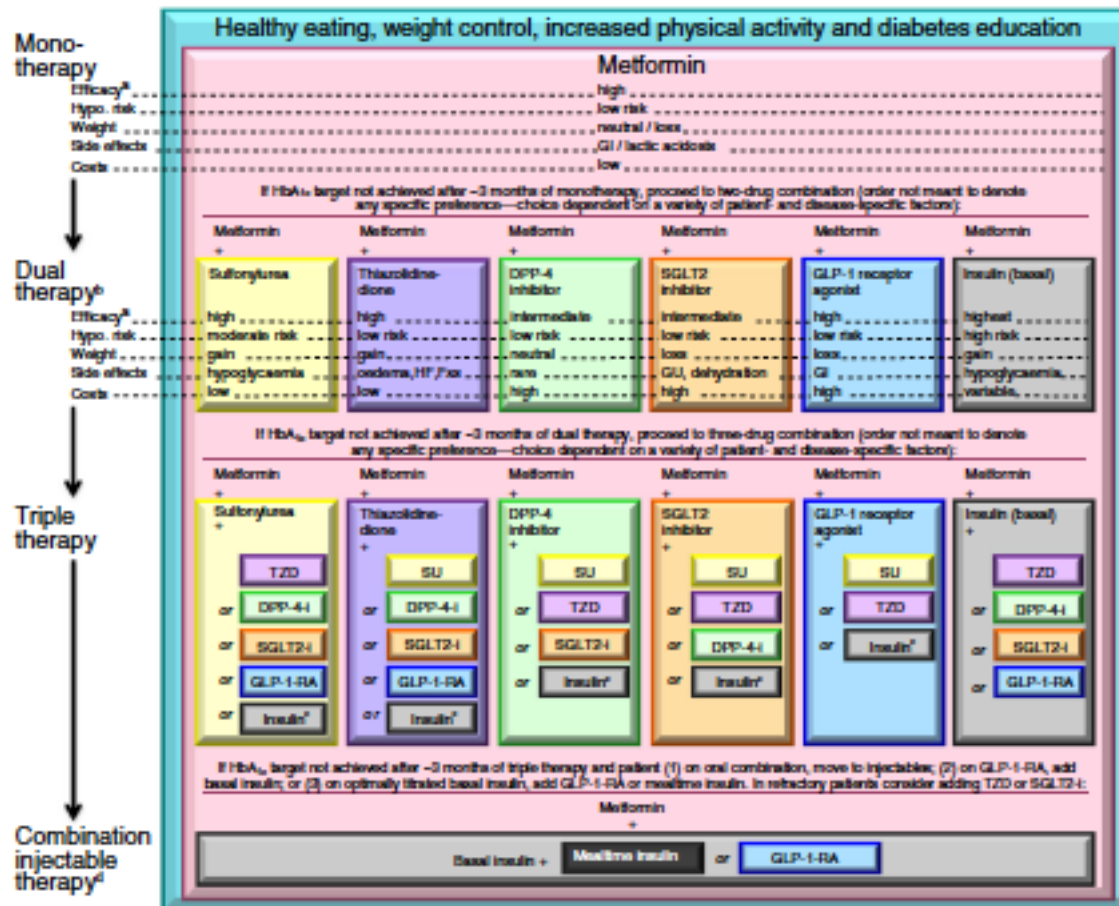
2015 Medikamente

| Properties of available glucose-lowering agents in the USA and Europe that may guide individualised treatment choices in patients with type 2 diabetes | | | | | | |
|--------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------|-------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------|
| Class | Compound(s) | Cellular mechanism(s) | Primary physiological action(s) | Advantages | Disadvantages | Cost ^e |
| Biguanides | • Metformin | Activates AMP-kinase (? other) | • ↓ Hepatic glucose production | • Extensive experience • No hypoglycaemia • ↓ CVD events (UKPDS) | • Gastrointestinal side effects (diarrhoea, abdominal cramping) • Lactic acidosis risk (rare) • Vitamin B ₁₂ deficiency • Multiple contraindications: CKD, acidosis, hypoxia, dehydration, etc. | Low |
| Sulfonylureas | 2nd generation • Glibenclamide/glyburide • Glipizide • Glimepiride • Gliclazide ^b | Closes K _{ATP} channels on beta cell plasma membranes | • ↑ Insulin secretion | • Extensive experience • ↓ Microvascular risk (UKPDS) | • Hypoglycaemia • ↑ Weight • ? Blunts myocardial ischaemic preconditioning • Low durability | Low |
| Meglitinides (glinides) | • Repaglinide • Nateglinide | Closes K _{ATP} channels on beta cell plasma membranes | • ↑ Insulin secretion | • ↓ Postprandial glucose excursions • Dosing flexibility | • Hypoglycaemia • ↑ Weight • ? Blunts myocardial ischaemic preconditioning • Frequent dosing schedule | Moderate |
| TZDs | • Pioglitazone ^c • Rosiglitazone ^d | Activates the nuclear transcription factor PPAR-γ | • ↑ Insulin sensitivity | • No hypoglycaemia • Durability • ↑ HDL-C • ↓ Triacylglycerols (pioglitazone) • ? ↓ CVD events (PROactive, | • ↑ Weight • Oedema/heart failure • Bone fractures • ↑ LDL-C (rosiglitazone) • ? ↑ MI (meta-analysis, rosiglitazone) | Low |
| Insulins | • Rapid-acting analogues – Lispro – Aspart – Glulisine • Short-acting – Human Regular • Intermediate-acting – Human NPH • Basal insulin analogues – Glargine – Detemir – Degludec ^b • Pre-mixed | Activates insulin receptors | • ↑ Glucose disposal • ↓ Hepatic glucose production • Other | • Nearly universal response • Theoretically unlimited efficacy • ↓ Microvascular risk (UKPDS) | • Training requirements • Hypoglycaemia • Weight gain • ? Mitogenic effects • Injectable • Training requirements • Patient reluctance | Variable ^e |

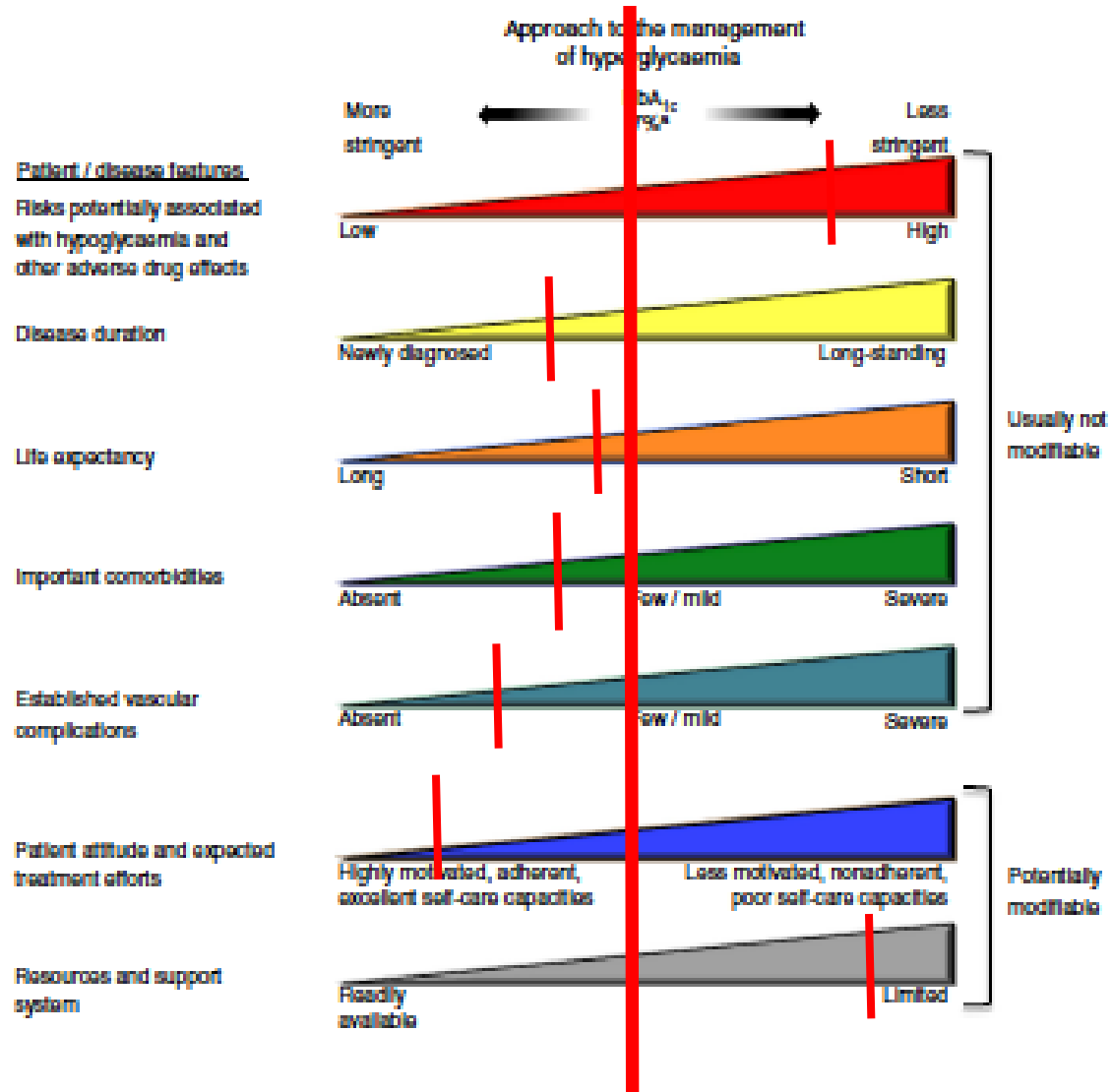
2015 Medikamente

| | | | | | | |
|-------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|
| DPP-4 inhibitors | <ul style="list-style-type: none"> • Sitagliptin • Vildagliptin^b • Saxagliptin • Linagliptin • Alogliptin | Inhibits DPP-4 activity, increasing postprandial active incretin (GLP-1, GIP) concentrations | <ul style="list-style-type: none"> • ↑ Insulin secretion (glucose-dependent) • ↓ Glucagon secretion (glucose-dependent) | <ul style="list-style-type: none"> • No hypoglycaemia • Well tolerated | <ul style="list-style-type: none"> • Angioedema/urticaria and other immuno-mediated dermatological effects • ? Acute pancreatitis • ? ↑ Heart failure hospitalisations • ? Rheinitis | High |
| SGLT2 inhibitors | <ul style="list-style-type: none"> • Canagliflozin • Dapagliflozin^c • Empagliflozin | Inhibits SGLT2 in the proximal nephron | <ul style="list-style-type: none"> • Blocks glucose reabsorption by the kidney, increasing glycosuria | <ul style="list-style-type: none"> • No hypoglycaemia • ↓ Weight • ↓ Blood pressure • Effective at all stages of T2DM | <ul style="list-style-type: none"> • Genitourinary infections • Polyuria • Volume depletion/hypotension/dizziness • ↑ LDL-C • ↑ Creatinine (transient) | High |
| GLP-1 receptor agonists | <ul style="list-style-type: none"> • Exenatide • Exenatide extended-release • Liraglutide • Albiglutide • Lixisenatide^b • Dulaglutide | Activates GLP-1 receptors | <ul style="list-style-type: none"> • ↑ Insulin secretion (glucose-dependent) • ↓ Glucagon secretion (glucose-dependent) • Slows gastric emptying • ↑ Satiety | <ul style="list-style-type: none"> • No hypoglycaemia • ↓ Weight • ↓ Postprandial glucose excursions • ↓ Some cardiovascular risk factors | <ul style="list-style-type: none"> • Gastrointestinal side effects (nausea/vomiting/diarrhoea) • ↑ Heart rate • ? Acute pancreatitis • C cell hyperplasia/medullary thyroid tumours in animals • Injectable • Training requirements | High |

2015 Behandlungsbaum ADA / EASD



Zeit der individualisierten Medizin



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Key Messages 2015

- viele Patienten jetzt und noch mehr in Zukunft
- viele Medikamente
- viel Geld im Spiel
- jeder Patient ist individuell und hat sein eigenes HbA1c Ziel
- noch wenig Erfahrungen mit den neuen Medikamenten
- nicht nur HbA1c denken
aber alle kardio-vaskuläre Risiken aggressiv behandeln

