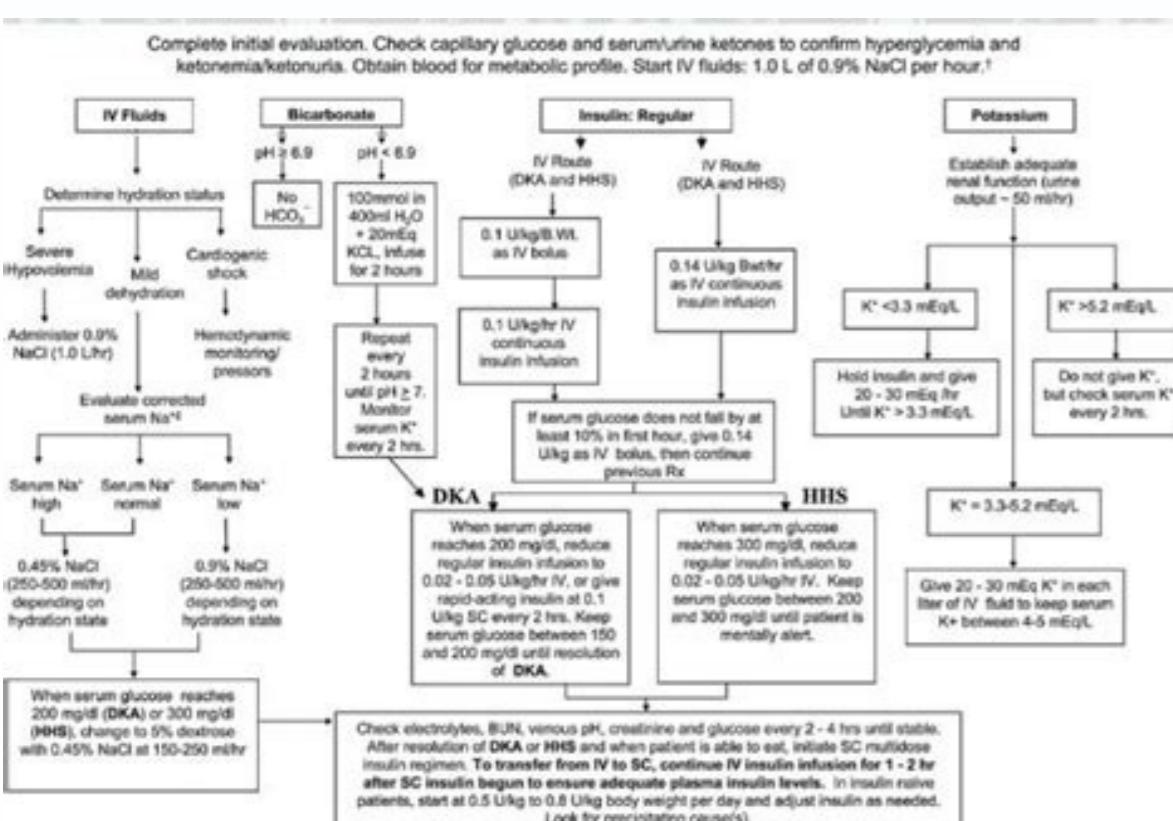


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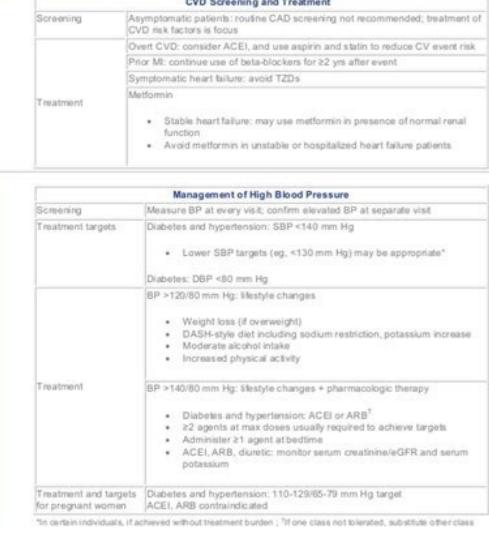
Kitabchi AE, Umpleby GE, Miles JM, et al. Hyperglycemic crises in adult patients with diabetes: a consensus statement from the American Diabetes Association. Diabetes Care. 2009;32:1345-1345.



American Diabetes Association (ADA) 2014 Guidelines  
Summary Recommendations from NDEI

Standards of medical care in diabetes—2014. Diabetes Care. 2014;37(suppl 1):S1-S80.

Refer to source document for full recommendations, including level of evidence rating.



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Source: American Diabetes Association.

Standards of medical care in diabetes—2014. Diabetes Care. 2014;37(suppl 1):S1-S80.

Refer to source document for full recommendations, including level of evidence rating.

ADA 2014 Guidelines

Criteria for Diabetes Diagnosis

A1C ≥ 6.5%  
Patient in lab using HbA1c method and standardized to DCCT assay

or

FPG ≥ 126 mg/dL (7.0 mmol/L)  
Fasting plasma glucose

or

2-hr PG ≥ 200 mg/dL (11.1 mmol/L) during OGTT (75 g)

or

Random PG ≥ 200 mg/dL (11.1 mmol/L)

In absence of unequivocal hyperglycemia, result to be confirmed by repeat testing.

FPG=fasting plasma glucose; OGTT=oral glucose tolerance test. Polymerase chain reaction (PCR)-based assays for A1C are not recommended.

Source: American Diabetes Association. Diabetes Care. 2014;37(suppl 1):S1-S80.

ADA 2014 Guidelines

Frequency of A1C Testing

Perform A1C test

At least 2 times each year in patients who are achieving treatment goals and have stable glycemic control

Quarterly in patients whose therapy has changed or who are not meeting glycemic targets

Point-of-care A1C testing allows for more timely treatment changes

American Diabetes Association. Diabetes Care. 2014;37(suppl 1):S1-S80.

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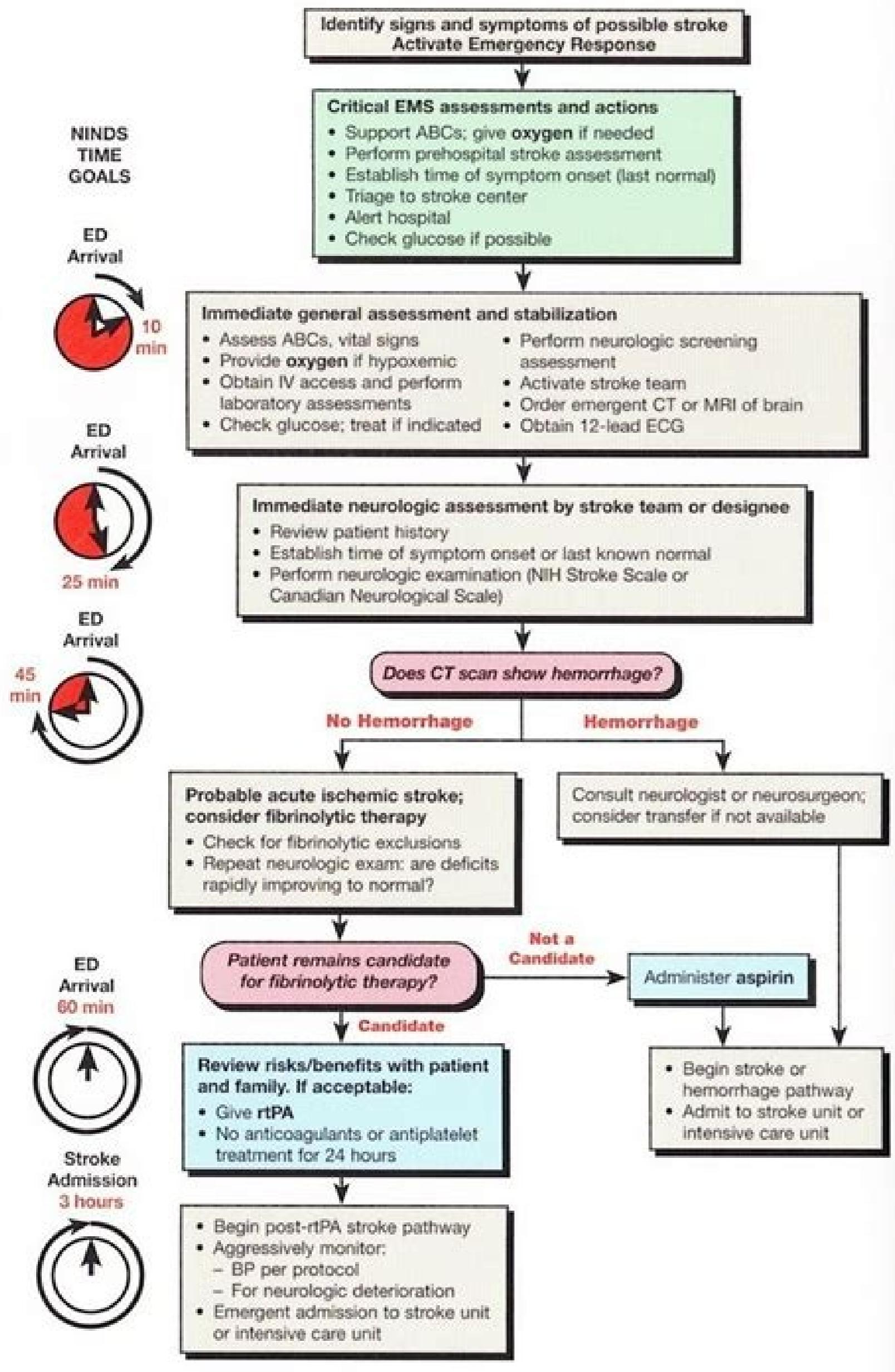
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## Suspected Stroke Algorithm: Goals for Management of Stroke



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Examples are Ozempic, Victoza, and Trulicity. Skip to Main Content Skip Nav Destination Skip to Main Content Skip Nav Destination The ADA guidelines still recommend Metformin as the first drug for patients with diabetes type 2 (DM2). In addition, the algorithm is formulated to be consistent with American Association of Clinical Endocrinologists (AACE) position statements on adiposity- and dysglycemia-based chronic disease models for early and sustainable preventive care. The above drug algorithmic guidance is general. The ADA recommends the following drugs as the third line option: GLP1 agonist if already Metformin + SGLT2 inhibitor and SGLT2 inhibitor if the patient is taking Metformin + GLP1 agonist. The recommendations are based on an extensive review of the clinical diabetes literature, supplemented with input from ADA staff and the medical community at large. The ultimate clinical decision is based on medication tolerability, cost, clinical setting, glucose control, comorbidities, and patient's preference. GTALSO SEE: Diabetes GuidelinesADA GuidelinesDiabetes Medication use in DM2Metformin is the preferred initial pharmacologic agent for the treatment of type 2 diabetes. Once initiated, metformin should be continued as long as it is tolerated and not contraindicated; other agents, including insulin, should be added to metformin. Early combination therapy can be considered in some patients at treatment initiation to extend the time to treatment failure. The early introduction of insulin should be considered if there is evidence of ongoing Catabolism (weight loss)/Symptoms of hyperglycemia/A1c levels >10%. Blood glucose ≥300 mg/dL A patient-centered approach should be used to guide the choice of pharmacologic agents. Such medications are Invokana, Jardiance, and Farxiga. In the accompanying algorithm, a chart summarizing the attributes of the antihyperglycemic class appear at the end. The advantages of Metformin are its effects in lowering A1c, being inexpensive, improving insulin resistance, potential weight loss, not causing hypoglycemia, and having the most extended clinical safety data - since its approval in 1957 in France and 1995 in the U.S. If the patient has established risk factors for cardiovascular disease, then a GLP1 agonist with proven CVD benefits is the recommended second-line medication. Consideration includes cardiovascular comorbidities, hypertension, and/or obesity. Examples of GLP1 agonists include exenatide, liraglutide, dulaglutide, semaglutide, and albiglutide. SGLT2 inhibitors include empagliflozin, canagliflozin, dapagliflozin, and ertugliflozin. DPP4 inhibitor include sitagliptin, linagliptin, and saxagliptin. TZD if heart failure is absent. In addition, the algorithm provides recommendations for blood pressure (BP) and lipid control, the two most important risk factors for atherosclerotic cardiovascular disease (ASCVD). This algorithm supplements the AACE and American College of Endocrinology (ACE) 2015 Clinical Practice Guidelines for Developing a Diabetes Mellitus Comprehensive Care Plan (6) and is organized into discrete sections that address the following topics: the founding principles of the algorithm, lifestyle therapy, obesity, prediabetes, management of hypertension and dyslipidemia, and glucose control with noninsulin antihyperglycemic agents and insulin. However, if a person with DM2 has heart failure or chronic kidney disease — defined by LVEF 300 — an SGLT2 inhibitor should be used. It is now clear that the progressive pancreatic beta-cell defect that drives the deterioration of metabolic control over time begins early and may be present before the diagnosis of T2D (1-3). Since originally drafted in 2013, the algorithm has been updated as new therapies, management approaches, and important clinical data have emerged. In addition to advocating glycemic control to reduce microvascular complications, this document highlights obesity and prediabetes as underlying risk factors for the development of T2D and associated macrovascular complications.

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