



COUNCIL OF THE DISTRICT OF COLUMBIA
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WASHINGTON, D.C. 20004

February 7, 2023

Nyasha Smith, Secretary
Council of the District of Columbia
1350 Pennsylvania Avenue, N.W.
Washington, DC 20004

Dear Secretary Smith,

Today, along with Councilmembers Charles Allen, Janeese Lewis George, Anita Bonds, Vincent C. Gray, Christina Henderson, Robert C. White, Jr., and Trayon White, Sr., I am introducing the Prior Authorization Reform Amendment Act of 2023. Please find enclosed a signed copy of the legislation.

In recent years, health insurers across the United States have adopted a new practice, wherein patients and their medical providers are required to seek prior authorization for certain medications, medical procedures, or other medical care. Prior authorization requirements mean that an insurer is able to overrule the treatment prescribed by a patient's medical provider—and make that determination without ever seeing the patient or their medical records. While insurers may claim that prior authorization is currently required for only complex procedures, a growing number of basic, everyday treatments require prior authorization; in fact, more and more, it seems that the cost of treatment, not a determination of medical necessity for different care, is driving what medications and procedures require prior authorization.

When unregulated, prior authorization requirements can and do cause meaningful harm to patients. Seeking a prior authorization (or appealing a denial) can take weeks or even months, during which time patients will typically go without care. This not only means the patient continues to suffer, but some conditions may worsen over time. Unfortunately, these lengthy delays inure to the benefit of insurers as patients, not wanting to wait for care, may seek different, possibly less effective treatment that doesn't require prior authorization. And, where that alternate treatment is less expensive, the insurer saves money.

Making matters worse, insurers may not make prior authorization requirements clear and accessible to patients. Some insurers may not include full information on prior authorization requirements on their website and decline to provide clarity via e-mail or the phone; often, patients may learn the grounds for a denial of a prior authorization and how to cure via letters sent snail mail, further delaying care for these patients. These delays and the lack of clarity on how one might successfully receive prior authorization for a treatment also push patients to seek cheaper, sometimes less effective care that does not require this approval. It is clear: unregulated prior

authorization requirements result in slower, worse quality care for patients, with the only benefits flowing to insurance companies.

In fact, prior authorization requirements also create problems for medical providers. Doctors report investing a growing number of staff hours to processing prior authorization requests or appeals on behalf of patients; that time is even greater in states, like the District, that do not regulate these processes, as insurers may use different forms, processes, and review standards. Some medical providers have even reported having to bring on additional staff to handle prior authorizations. This means higher overhead for doctor's offices—costs that ultimately get passed on to patients. Dealing with these processes also mean medical providers have less time to care for patients.

More than forty states have passed legislation or adopted regulations to address insurer prior authorization practices, and this legislation would bring the District in line with those jurisdictions. The bill sets explicit, reasonable timelines for insurers to respond to prior authorization requests and appeals and lays out the qualifications of personnel qualified to make these determinations (as some insurers have personnel without appropriate subject matter training reviewing prior authorization requests). The legislation would also clarify how insurers are to make information on prior authorization determinations available to patients and their medical providers and require that insurers accept and use the electronic NCPDP SCRIPT Standard ePA transaction, the recommended, standardized method for submission and review of prior authorization requests. Finally, and perhaps most importantly, the bill would prohibit insurers from requiring prior authorization for a treatment based solely on cost. Separately, this legislation would make a small change to require that employers provide timely notice to employees of medications and treatments covered under their insurer's standard health benefit plan, but not covered under the negotiated terms of the employer's bespoke plan; this language will help ensure employees have full knowledge of what is and what isn't covered under various employer health benefit plans, and can make a fully educated choice about which coverage to choose.

This legislation is being introduced with the input and support of the Medical Society of the District of Columbia. As noted, more than forty states have already acted to regulate this practice, recognizing the meaningful impact needless delays and denials of care due to prior authorization practices have on patients' health and wellbeing; of note, those states have not seen meaningful changes in the cost of medications or care due to regulation this practice. Importantly, with this legislation, the District will help ensure patients do not face unnecessarily barriers to timely, medically appropriate care.

Should you have any questions about this legislation, please contact my Legislative Counsel, Kristin Ewing, at kewing@dccouncil.gov.

Thank you,



Brooke Pinto



Councilmember Charles Allen



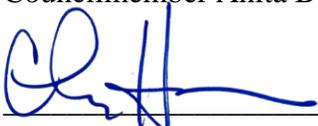
Councilmember Brooke Pinto



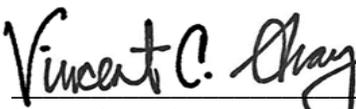
Councilmember Anita Bonds



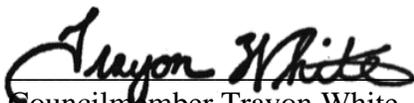
Councilmember Janeese Lewis George



Councilmember Christina Henderson



Councilmember Vincent C. Gray



Councilmember Trayon White, Sr.



Councilmember Robert C. White, Jr.

A BILL

IN THE COUNCIL OF THE DISTRICT OF COLUMBIA

To prescribe the manner in which a utilization review entity is to make available information on prior authorization requirements and restrictions; to set notice requirements for prior authorization determinations; to lay out the minimum length that a prior authorization is to be considered valid, to set the qualifications for personnel authorized to make adverse determinations; to permit enrollees to appeal an adverse determination and to set deadlines for submissions of appeals; to set qualifications for personnel authorized to review appeals of adverse determinations; to prescribe utilization review entities' obligations in terms of reviewing requests for prior authorization for non-urgent, urgent, and emergency; to permit utilization review entities to require prior authorization only based on a determination of medical necessity for different care and to prohibit a utilization review entity from requiring prior authorization for a treatment solely based on cost; to prohibit a utilization review entity from revoking, limiting, condition, or restricting a prior authorization if care was provided within 45 days of receipt of the prior authorization; to require that a utilization review entity honor a prior authorization granted by a previous utilization review entity for at least the first 60 days of coverage; to clarify that health services are to be deemed authorized if a utilization review entity fails to comply with this act; and to require utilization review entities using prior authorization to make certain statistics available to the public; to amend the Uniform Health Insurance Claims Forms Act of 1995 to require that, by January 1, 2024, all utilization review entities accept and respond to prior authorization requests using the NCPDP SCRIPT Standard ePA transaction; and to amend the Health Insurance Portability and

31 Accountability Federal Law Conformity and No-Fault Motor Vehicle Insurance Act of
32 1998 to require that employers provide notice of employees of treatments, including
33 particular services or medications, not included in a negotiated health benefit plan but
34 including in the standard health benefit plan or formulary offered by the health insurer.
35

36 BE IT ENACTED BY THE COUNCIL OF THE DISTRICT OF COLUMBIA, That this
37 act may be cited as the “Prior Authorization Reform Amendment Act of 2023”.

38 Sec. 2. Definitions.

39 (a) “Adverse determination” means a decision by a utilization review entity that the
40 health care services furnished or proposed to be furnished to an enrollee are not medically
41 necessary or are experimental or investigational; and benefit coverage is therefore denied,
42 reduced, or terminated.

43 (b) “Emergency health care services” means those health care services that are provided
44 in an emergency facility after the sudden onset of a medical condition that manifests itself by
45 symptoms of sufficient severity, including severe pain, that the absence of immediate medical
46 attention could reasonably be expected by a prudent layperson, who possesses an average
47 knowledge of health and medicine, to result in placing the patient's health in serious jeopardy,
48 serious impairment to bodily function, or serious dysfunction of any bodily organ or part.

49 (c) “Enrollee” means an individual eligible to receive health care benefits by a health
50 insurer pursuant to a health plan or other health insurance coverage. The term “enrollee”
51 includes an enrollee’s legally authorized representative.

52 (d) “Medication assisted treatment” means the use of medications, commonly in
53 combination with counseling and behavioral therapies, to provide a comprehensive approach to
54 the treatment of substance use disorders.

55 (e) “Prior authorization” means the process by which utilization review entities determine
56 the medical necessity or medical appropriateness of covered health care services prior to the
57 rendering of such health care services. “Prior authorization” also includes any health insurer or
58 utilization review entity’s requirement that an enrollee or health care provider notify the health
59 insurer or utilization review entity prior to providing a health care service.

60 (f) “Urgent health care service” means:

61 (1) A health care service that, in the opinion of a physician with knowledge of the
62 enrollee’s medical condition, if not receiving an expedited prior authorization:

63 (A) Could seriously jeopardize the life or health of the enrollee or the
64 ability of the enrollee to regain maximum function; or

65 (B) Could subject the enrollee to severe pain that cannot be adequately
66 managed without the care or treatment that is the subject of the utilization review; or

67 (2) Medication assisted treatment.

68 (g) “Utilization review entity” means an individual or entity that performs prior
69 authorization for one or more of the following entities:

70 (i) An employer with employees in the District;

71 (ii) An insurer that writes health insurance policies;

72 (iii) A preferred provider organization, or health maintenance organization; and

73 (iv) Any other individual or entity that provides, offers to provide, or administers
74 hospital, outpatient, medical, prescription drug, or other health benefits to a person treated by a
75 health care provider in the District under a policy, plan, or contract.

76 Sec. 3. Prior Authorization Requirements and Restrictions.

77 (a)(1) A utilization review entity shall make available any current prior authorization
78 requirements and restrictions, including formulary, (“prior authorization requirements”) in at
79 least the following ways:

80 (A) Posting of the prior authorization requirements on its website, in a
81 manner accessible to enrollees, health care providers, and the general public and without an
82 account;

83 (B) Emailing or providing a hard copy of the prior authorization
84 requirements to enrollees and health care providers upon request by telephone or in writing,
85 including a request via email; and

86 (C) Providing information on prior authorization requirements, upon
87 request, to enrollees or health care providers over the telephone.

88 (b) Prior authorization requirements shall:

89 (1) Be described in detail and easily understandable language;

90 (2) Include any written clinical criteria;

91 (3) Include a comprehensive listing of all drugs that require a prior authorization;

92 and

93 (4) Include the process for submitting and standards for considering, including
94 evidence-based guidelines, where possible, requests for:

95 (i) A prior authorization;

96 (ii) A reauthorization of a prior authorization after a previous prior
97 authorization has expired, and

98 (iii) Appeals of an adverse determination.

99 (c)(1) If a utilization review entity intends either to amend or replace the prior
100 authorization requirements, the changes shall not be deemed effective until the utilization review
101 entity's website has been updated to reflect the new language.

102 Sec. 4. Prior Authorization Determinations.

103 (a)(1) Where a utilization review entity makes a determination to grant or deny a prior
104 authorization, the enrollee and the health care provider submitting the request for a prior
105 authorization must be provided with notice within 24 hours of the determination.

106 (2) Notice provided under this section must include:

107 (A) The name and qualifications, pursuant to Section 6 of this Act, of the
108 personnel making the determination; and

109 (B) For an adverse determination:

110 (i) The grounds under the prior authorization requirements for
111 denying the prior authorization; and

112 (ii) Information on the enrollee's right to appeal, the process to file
113 an appeal, and a listing of information necessary to support an appeal of the adverse
114 determination.

115 (c)(1) A utilization review entity shall make information available on its website to
116 enrollees and the enrollee's health care provider on active requests for prior authorization and
117 requests made, at a minimum, to that utilization review entity in the preceding five years, and
118 shall include:

119 (A) A copy of any information or materials submitted by the enrollee's
120 health care provider to request or support a request for a prior authorization or reauthorization, or
121 appeal an adverse determination; the information or materials shall clearly show the date of any

122 submissions by the health care provider, the health care service prescribed by the health care
123 provider, and the basis, if any, provided by the health care provider for the health care service;
124 and

125 (B) A copy of notices of determination provided to the enrollee and health
126 care provider pursuant to subsection (a) of this section;

127 (b) Upon request of the enrollee or health care provider, a utilization review entity shall
128 make information on an adverse determination available via telephone, including the basis under
129 the prior authorization requirements for denying the prior authorization, information on the
130 enrollee's right to appeal, the process to file an appeal, and any information necessary to support
131 a successful appeal.

132 Sec. 5. Length of prior authorization.

133 (a) Except for as provided in subsection (b) of this section, a prior authorization for shall
134 be valid for at least one year from the date the health care provider receives the prior
135 authorization. The prior authorization shall remain valid regardless of any changes in dosage for
136 a prescription drug prescribed by the health care provider; provided, that utilization review
137 entities may rescind prior authorization for dosages exceeding limitations set in federal or
138 District law or regulations.

139 (b) If a utilization review entity requires a prior authorization for a health care service for
140 the treatment of a chronic or long-term care condition, the prior authorization shall remain valid
141 for the length of the treatment and the utilization review entity may not require the enrollee to re-
142 obtain a prior authorization for the health care service.

143 Sec. 6. Personnel qualified to make adverse determinations.

144 (c) A utilization review entity must ensure that all adverse determinations are made by a
145 physician who:

146 (1) Possesses a current and valid non-restricted license to practice medicine in the
147 District of Columbia;

148 (2) Is of the same specialty as a physician who typically manages the medical
149 condition or disease or provides the health care service involved in the request;

150 (3) Makes the adverse determination under the clinical direction of one of the
151 utilization review entity's medical directors who is responsible for the provision of health care
152 services provided to enrollees in the District of Columbia, and who is licensed in the District of
153 Columbia.

154 Sec. 7. Consultation prior to issuing an adverse determination

155 (a) If a utilization review entity is questioning the medical necessity of a health care
156 service, the utilization review entity must notify the enrollee's health care provider that medical
157 necessity is being questioned. Prior to issuing an adverse determination, the enrollee's health
158 care provider must have the opportunity to discuss the medical necessity of the health care
159 service on the telephone with the physician who will be responsible for determining
160 authorization of the health care service under review.

161 Sec. 8. Appeals.

162 (a)(1) A utilization review entity shall allow an enrollee to appeal an adverse
163 determination. Any appeal submitted within 15 calendar days of the enrollee's receipt of notice
164 of the adverse determination shall be treated as timely.

165 (2) A utilization review entity shall permit an appeal to be submitted at least via
166 its website or in hard copy.

167 (3) Appeals submitted in hard copy shall be considered timely where the appeal is
168 postmarked within 15 calendar days of the enrollee's receipt of notice of the adverse
169 determination.

170 (b)(1) The enrollee and the health care provider submitting the original request for a prior
171 authorization must be provided notice within 24 hours of a determination on an appeal of an
172 adverse determination.

173 (2) Notice provided under this subsection must include:

174 (A) The name and qualifications, pursuant to Section 9 of this Act, of the
175 physician reviewing the appeal; and

176 (B) The grounds under the prior authorization requirements for the
177 physician's determination.

178 Sec. 9. Personnel qualified to review appeals.

179 (a) A utilization entity must ensure that all appeals are reviewed by a physician. The
180 physician must:

181 (1) Possess a current and valid non-restricted license to practice medicine in the
182 District;

183 (2) Be in active practice in the same specialty as a physician who typically
184 manages the medical condition or disease and have practiced that specialty for at least 5 years;

185 (3) Be knowledgeable of, and have experience providing, the health care services
186 under appeal;

187 (4) Not be employed by a utilization review entity or be under contract with the
188 utilization review entity other than to participate in one or more of the utilization review entity's

189 health care provider networks or to perform reviews of appeals, or otherwise have any financial
190 interest in the outcome of the appeal;

191 (5) Not have been directly involved in making the adverse determination; and

192 (b) In reviewing an appeal, the physician must consider all known clinical aspects of the
193 health care service under review, including but not limited to, a review of all pertinent medical
194 records provided to the utilization review entity by the enrollee's health care provider, any
195 relevant records provided to the utilization review entity by a health care facility, and any
196 medical literature provided to the utilization review entity by the health care provider.

197 Sec. 10. Utilization review entities' obligations with respect to prior authorizations in
198 non-urgent, urgent, and emergency circumstances.

199 (a) If a utilization review entity requires prior authorization of a health care service, the
200 utilization review entity must grant the prior authorization or make an adverse determination and
201 notify the enrollee and the enrollee's health care provider of the prior authorization or adverse
202 determination within 3 business days of obtaining all information required; if the determination
203 is not made within that time frame, such services shall be deemed approved.

204 (b) A utilization review entity must grant a prior authorization or make an adverse
205 determination concerning urgent care services and notify the enrollee and the enrollee's health
206 care provider of that determination, not later than 24 hours after receiving all information
207 required if the determination is not made within that time frame, such services shall be deemed
208 approved.

209 (c)(1) A utilization review entity cannot require prior authorization for pre-hospital
210 transportation or for the provision of emergency health care services, including emergency health
211 care services to screen and stabilize an enrollee.

212 (2) A utilization review entity shall allow an enrollee and the enrollee's health
213 care provider a minimum of 24 hours following an emergency admission or provision of
214 emergency health care services for the enrollee or health care provider to notify the utilization
215 review entity of the admission or provision of health care services. If the admission or health
216 care service occurs on a holiday or weekend, a utilization review entity cannot require
217 notification until the next business day after the admission or provision of the health care
218 services.

219 (3) If a health care provider certifies in writing to a utilization review entity within
220 72 hours of an enrollee's admission that the enrollee's condition required emergency health care
221 services, that certification will create a presumption that the emergency health care services were
222 medically necessary and such presumption may be rebutted only if the utilization review entity
223 can establish, with clear and convincing evidence, that the emergency health care services were
224 not medically necessary.

225 (4) The medical necessity or appropriateness of emergency health care services
226 cannot be based on whether those services were provided by participating or nonparticipating
227 providers. Restrictions on coverage of emergency health care services provided by
228 nonparticipating providers cannot be greater than restrictions that apply when those services are
229 provided by participating providers.

230 (d) For purposes of this section, "required information" includes the results of any face-
231 to-face clinical evaluation or second opinion that may be required.

232 Sec. 11. Prior Authorization Limitations.

233 (a) A utilization review entity may only require prior authorization for a health care
234 service based on a determination of medical necessity for different care or that the proposed care

235 is experimental or investigational in nature. A utilization review entity may not require prior
236 authorization solely based on the cost of a health care service.

237 (b) A utilization review entity may not require prior authorization for the provision of
238 medication-assisted treatment for the treatment of opioid-use disorder.

239 Sec. 12. Retrospective denial.

240 The utilization review entity may not revoke, limit, condition, or restrict a prior
241 authorization if care is provided within 45 working days from the date the health care provider
242 received the prior authorization.

243 Sec. 13. Continuity of care for enrollees.

244 (a) A utilization review entity shall honor a prior authorization granted to an enrollee
245 from a previous utilization review entity for at least the initial 60 days of an enrollee's coverage
246 under a new health plan; provided, that the utilization review entity may condition honoring the
247 prior authorization on receipt of information documenting the previous utilization review entity's
248 grant of prior authorization.

249 (b) During the time period described in paragraph (a) of this subsection, a utilization
250 review entity may perform its own review to grant a prior authorization.

251 (c) If there is a change in coverage of, or approval criteria for, a previously authorized
252 health care service, the change in coverage or approval criteria shall not apply to an enrollee who
253 received prior authorization before the effective date of the change for the length of the prior
254 authorization's eligibility.

255 (d) A utilization review entity shall continue to honor a prior authorization it has granted
256 to an enrollee when the enrollee changes products under the same health insurance company.

257 Sec. 14. Health care services deemed authorized if a utilization review entity fails to
258 comply with the requirements of this Act.

259 Any failure by a utilization review entity to comply with the deadlines and other
260 requirements specified in this Act shall result in the health care services in question to be deemed
261 authorized by the utilization review entity.

262 Sec. 15. Data Collection.

263 Utilization review entities using prior authorization shall make statistics available
264 regarding prior authorizations, adverse determinations, and appeals on their website in a readily
265 accessible format. They should include categories for approvals, adverse determinations, and
266 appeals broken down by:

- 267 (1) Specialty of physician reviewing the request for prior authorization or appeal;
- 268 (2) Type of medication, test, procedure, or treatment (“health care service”);
- 269 (3) Indication offered;
- 270 (4) Reason for denial;
- 271 (5) If appealed;
- 272 (6) If approved or denied on appeal;
- 273 (7) The time between submission of the request for prior authorization and the
274 utilization review entity’s determination; and
- 275 (8) The time between submission of an appeal of an adverse determination and
276 the utilization review entity’s determination.

277 Sec. 16. The Uniform Health Insurance Claim Forms Act of 1995, effective February 27,
278 1996 (D.C. Law 11-89; D.C. Code §31-3201) is amended by adding a new subsection (c) to read
279 as follows:

280 “(c)(1) No later than January 1, 2024, a utilization review entity must accept and respond
281 to prior authorization requests under the pharmacy benefit through a secure electronic
282 transmission using the NCPDP SCRIPT Standard ePA transactions. Facsimile, propriety payer
283 portals, electronic forms, or any other technology not directly integrated with a physician’s
284 electronic health record/electronic prescribing system shall not be considered secure electronic
285 transmission.

286 “(2) For the purposes of this subsection:

287 “(A) “NCPDP SCRIPT Standard” means the National Council for
288 Prescription Drug Programs SCRIPT Standard Version 2013101, or the most recent standard
289 adopted by the United States Department of Health and Human Services.

290 “(B) “Prior authorization” means the process by which utilization review
291 entities determine the medical necessity or medical appropriateness of covered health care
292 services prior to the rendering of such health care services. “Prior authorization” also includes
293 any health insurer or utilization review entity’s requirement that an enrollee or health care
294 provider notify the health insurer or utilization review entity prior to providing a health care
295 service.

296 “(C) “Utilization review entity” means an individual or entity that
297 performs prior authorization for one or more of the following entities:

298 “(i) An employer with employees in the District;

299 “(ii) An insurer that writes health insurance policies;

300 “(iii) a preferred provider organization, or health maintenance

301 organization; and

302 “(iv) any other individual or entity that provides, offers to provide,
303 or administers hospital, outpatient, medical, prescription drug, or other health benefits to a
304 person treated by a health care provider in the District under a policy, plan, or contract.”

305 Sec. 17. The Health Insurance Portability and Accountability Federal Law Conformity
306 and No-Fault Motor Vehicle Insurance Act of 1998, effective April 13, 1999 (D.C. Law 12-209;
307 D.C. Official Code § 31-3301.01), is amended by adding a new Section 313e to read as follows:

308 “Section 313e. Negotiated health benefit plans.

309 “(a) Where an employer negotiates an employee health benefit plan with a health insurer
310 such that treatment, including particular services or medications, covered under the negotiated
311 health benefit plan offered to employees differs from the standard health benefit plan or
312 formulary offered by the health insurer, the employer shall provide notice to all employees,
313 regardless of whether they are enrolled in the negotiated health benefit plan, of any treatments,
314 including particular services or medications, covered under the standard health benefit plan or
315 formulary but not covered under the negotiated health benefit plan or formulary offered to
316 employees.

317 “(b) Notice under subsection (a) of this section shall be provided to employees:

318 “(1) At least 30 days prior to the conclusion of any open enrollment period; and

319 “(2) Within 30 days after the employer and health insurer finalize terms of
320 coverage under a negotiated health benefit plan.

321 “(c) For the purposes of

322 Sec. 18. Fiscal impact statement.

323 The Council adopts the fiscal impact statement in the committee report as the fiscal
324 impact statement required by section 4a of the General Legislative Procedures Act of 1975,
325 approved October 16, 2006 (120 Stat. 2038; D.C. Official Code § 1-301.47a).

326 Sec. 19. Effective date.

327 This act shall take effect following approval by the Mayor (or in the event of veto by the
328 Mayor, action by the Council to override the veto), a 30-day period of congressional review as
329 provided in section 602(c)(1) of the District of Columbia Home Rule Act, approved December
330 24, 1973 (87 Stat. 813; D.C. Official Code § 1-206.02(c)(1)), and publication in the District of
331 Columbia Register.